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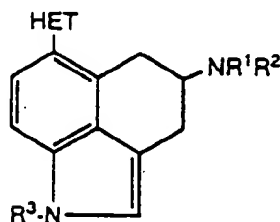
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(54) **6-Heterocyclyl-4-amino-1,3,4,5-tetrahydrobenz CD Indoles for treating motion sickness and vomiting.**

(57) The use of a compound, or pharmaceutically acceptable salt thereof, of the formula



wherein :

R¹ is hydrogen, C₁-C₄ alkyl, C₃-C₄ alkenyl, cyclopropylmethyl, aryl(C₁-C₄ alkyl), -(CH₂)_nS(C₁-C₄ alkyl), -C(O)R⁴, or -(CH₂)_nC(O)NR⁵R⁶ ;

R² is hydrogen, C₁-C₄ alkyl, cyclopropylmethyl or C₃-C₄ alkenyl ;

R³ is hydrogen, C₁-C₄ alkyl or an amino blocking group ;

n is 1-4 ;

R⁴ is hydrogen, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy or phenyl ;

R⁵ and R⁶ are independently hydrogen, C₁-C₄ alkyl, or C₅-C₈ cycloalkyl with the proviso that when one of R⁵ or R⁶ is a cycloalkyl the other is hydrogen ;

HET is a tetrazolyl ring, a substituted tetrazolyl ring or an aromatic 5- or 6-membered heterocyclic ring, said ring having from one to three heteroatoms which are the same or different and which are selected from the group consisting of sulfur, oxygen, and nitrogen with the proviso that the 6-membered heterocyclic ring can only contain carbon and nitrogen and with the further proviso that the 5-membered ring contains no more than one oxygen or one sulfur but not both oxygen and sulfur ; to prepare a medicament for treating emesis or motion sickness ; and a compound of the formula above wherein HET is an optionally substituted tetrazolyl ring.

The present invention relates to a method of treating emesis and motion sickness and pharmaceutical formulations suitable therefor.

Flaugh in U.S. Patent No. 4,576,959 (issued 1986) disclosed a family of 6-substituted-4-dialkylamino-1,3,4,5-tetrahydrobenz[cd]indoles which are described as central serotonin agonists. Leander in U.S. Patent 4,745,126 (1988) disclosed a method for treating anxiety in humans employing a 4-substituted-1,3,4,5-tetrahydrobenz[cd]indole-6-carboxamide derivative.

European Patent Application 399,982 discloses certain heterocyclic-substituted aminotetralins. These compounds are disclosed as being serotonin agonists, partial agonists or antagonists.

Despite the progress of science as represented above, many mammals, both humans and animals, continue to be afflicted with emesis and motion sickness. Accordingly, the need continues for safer, more selective, drugs which can be used to treat such diseases. As such, it is an object of the present invention to provide a novel method for treating emesis and motion sickness. Further objects of the present invention are to provide novel formulations suitable for use in the instantly claimed method, as well as novel compounds which can be used in such method.

The present invention provides a method of treating emesis or motion sickness in mammals comprising administering to a mammal in need of such treatment an effective dose of a compound, or pharmaceutically acceptable salt thereof, of the Formula 1



wherein:

R¹ is hydrogen, C₁-C₄ alkyl, C₃-C₄ alkenyl, cyclopropylmethyl, aryl (C₁-C₄ alkyl), -(CH₂)ₙS(C₁-C₄ alkyl), -C(O)R⁴, or -(CH₂)ₙC(O)NR⁵R⁶;

R² is hydrogen, C₁-C₄ alkyl, cyclopropylmethyl or C₃-C₄ alkenyl;

R³ is hydrogen, C₁-C₄ alkyl or an amino-blocking group;

n is 1-4;

R⁴ is hydrogen, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy or phenyl;

R⁵ and R⁶ are independently hydrogen, C₁-C₄ alkyl, or C₅-C₈ cycloalkyl, with the proviso that when one of R⁵ or R⁶ is a cycloalkyl the other is hydrogen;

HET is a tetrazolyl ring, a substituted tetrazolyl ring or an aromatic 5- or 6-membered heterocyclic ring, said ring having from one to three heteroatoms which are the same or different and which are selected from the group consisting of sulfur, oxygen, and nitrogen with the proviso that the 6-membered heterocyclic ring can only contain carbon and nitrogen and with the further proviso that a 5-membered ring may contain no more than one oxygen or one sulfur but not both oxygen and sulfur.

Compounds of Formula 1 have not heretofore been used to treat emesis or motion sickness. Accordingly, another embodiment of this invention is a pharmaceutical formulation adapted for the treatment of emesis or motion sickness comprising a compound of Formula 1, or a pharmaceutically acceptable salt thereof, in combination with one or more pharmaceutically acceptable carriers, diluents or excipients therefor.

Certain compounds of Formula 1, i.e., those compounds wherein HET is a tetrazolyl ring or a substituted tetrazolyl ring and R¹, R² and R³ are as defined for Formula 1, are novel. Accordingly, yet another embodiment of the present invention provides such novel compounds, processes for preparing and methods of using same.

As used herein, the term "alkyl" represents a straight or branched alkyl chain having the indicated number of carbon atoms. For example, "C₁-C₄ alkyl" groups are methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, *sec*-butyl, isobutyl and *tert*-butyl. "C₁-C₈ alkyl" groups include those listed for C₁-C₄ alkyl as well as *n*-pentyl, 2-methylbutyl, 3-methylbutyl, *n*-hexyl, 4-methylpentyl, *n*-heptyl, 3-ethylpentyl, 2-methylhexyl, 2,3-dimethylpentyl, *n*-octyl, 3-propylpentyl, 6-methylheptyl, and the like.

The term "C₃-C₄ alkenyl" refers to olefinically unsaturated alkyl groups such as -CH₂CH=CH₂, -CH₂CH₂CH=CH₂, -CH(CH₃)CH=CH₂ and the like.

The term "aryl" means an aromatic carbocyclic structure having six to ten carbon atoms. Examples of such ring structures are phenyl, naphthyl, and the like.

The term "cycloalkyl" means an aliphatic carbocyclic structure having the indicated number of carbon

atoms in the ring. For example, the term "C₃-C₇ cycloalkyl" means cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, and cycloheptyl.

The term "aryl (C₁-C₄ alkyl)" means an aryl structure joined to a C₁-C₄ alkyl group. Examples of such groups are benzyl, phenylethyl, α -methylbenzyl, 3-phenylpropyl, α -naphthylmethyl, β -naphthylmethyl, 4-phenylbutyl, and the like. Similarly the term "aryl (C₁-C₃ alkyl)" means an aromatic carbocyclic structure joined to a C₁-C₃ alkyl.

The C₁-C₈ alkyl, aryl, aryl (C₁-C₄ alkyl) and aryl (C₁-C₃ alkyl) groups can be substituted by one or two moieties. Typical aryl and/or alkyl substituents are C₁-C₃ alkoxy, halo, hydroxy, C₁-C₃ thioalkyl, nitro, and the like. Moreover, the aryl, aryl (C₁-C₄ alkyl) and aryl (C₁-C₃ alkyl) groups can also be substituted by a C₁-C₃ alkyl or a trifluoromethyl group.

In the foregoing, the term "C₁-C₃ alkyl" means any of methyl, ethyl, *n*-propyl, and isopropyl; the term "C₁-C₃ alkoxy" means any of methoxy, ethoxy, *n*-propoxy, and isopropoxy; the term "halo" means any of fluoro, chloro, bromo, and iodo; and the term "C₁-C₃ thioalkyl" means any of methylthio, ethylthio, *n*-propylthio, and isopropylthio.

Examples of substituted C₁-C₈ alkyl are methoxymethyl, trifluoromethyl, 6-chlorohexyl, 2-bromopropyl, 2-ethoxy-4-iodobutyl, 3-hydroxypentyl, methylthiomethyl, and the like.

Examples of substituted aryl are *p*-bromophenyl, *m*-iodophenyl, *p*-tolyl, *o*-hydroxyphenyl, β -(4-hydroxy)-naphthyl, *p*-(methylthio)phenyl, *m*-trifluoromethylphenyl, 2-chloro-4-methoxyphenyl, α -(5-chloro)naphthyl, and the like.

Examples of substituted aryl (C₁-C₄ alkyl) are *p*-chlorobenzyl, *o*-methoxybenzyl, *m*-(methylthio)- α -methylbenzyl, 3-(4'-trifluoromethylphenyl)propyl, *o*-iodobenzyl, *p*-methylbenzyl, and the like.

The term "amino-blocking group" is used herein as it is frequently used in synthetic organic chemistry, to refer to a group which will prevent an amino group from participating in a reaction carried out on some other functional group of the molecule, but which can be removed from the amine when it is desired to do so. Such groups are discussed by T. W. Greene in chapter 7 of Protective Groups in Organic Synthesis, John Wiley and Sons, New York, 1981, and by J. W. Barton in chapter 2 of Protective Groups in Organic Chemistry, J. F. W. McOmie, ed., Plenum Press, New York, 1973, which are incorporated herein by reference in their entirety. Examples of such groups include benzyl and substituted benzyl such as 3,4-dimethoxybenzyl, *o*-nitrobenzyl, and triphenylmethyl; those of the formula -COOR where R includes such groups as methyl, ethyl, propyl, isopropyl, 2,2,2-trichloroethyl, 1-methyl-1-phenylethyl, isobutyl, *t*-butyl, *t*-amyl, vinyl, allyl, phenyl, benzyl, *p*-nitrobenzyl, *o*-nitrobenzyl, and 2,4-dichlorobenzyl; acyl groups and substituted acyl such as formyl, acetyl, chloroacetyl, dichloroacetyl, trichloroacetyl, trifluoroacetyl, benzoyl, and *p*-methoxybenzoyl; and other groups such as methanesulfonyl, *p*-toluenesulfonyl, *p*-bromobenzenesulfonyl, *p*-nitrophenylethyl, and *p*-toluenesulfonylamino-carbonyl. Preferred amino-blocking groups are benzyl (-CH₂C₆H₅), acyl [C(O)R] or SiR₃ where R is C₁-C₄ alkyl, halomethyl, or 2-halo-substituted-(C₂-C₄ alkoxy).

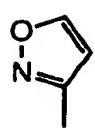
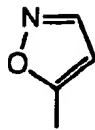
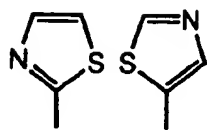
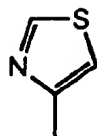
The term "aromatic 5- or 6-membered heterocyclic ring" refers to a ring containing from one to three heteroatoms which can be nitrogen, oxygen or sulfur. The 5-membered heterocyclic rings can contain carbon and nitrogen atoms and up to one oxygen or one sulfur but not one of each. In 5-membered rings not containing oxygen or sulfur, one nitrogen can be substituted with either a hydrogen, C₁-C₃ alkyl, phenyl or (C₁-C₃ alkyl)phenyl group. The 6-membered heterocyclic rings can contain carbon and nitrogen atoms only. The 5- or 6-membered rings can have one or two of the carbon atoms in the ring substituted independently with C₁-C₃ alkyl, halogen, OH, C₁-C₃ alkoxy, C₁-C₃ alkylthio, NH₂, CN or phenyl. Adjacent carbons in the heterocyclic ring may be connected with a -CH=CH-CH=CH- bridge to form a benzo-fused ring on the heterocycle.

These aromatic 5- or 6-membered heterocyclic rings can be either substituted or unsubstituted and include furan, thiophene, thiazole, oxazole, isoxazole, isothiazole, oxadiazole, thiadiazole, pyridine, pyridazine, pyrimidine, pyrazine, pyrrole, pyrazole, imidazole, and triazole. The heterocyclic ring can be attached to the benzene ring by any carbon in the heterocyclic ring, for example, 2- or 3-furan.

The term "substituted tetrazolyl ring" refers to a tetrazolyl ring system which has a C₁-C₃ alkyl or phenyl substituent on the 2-position nitrogen atom of such ring system.

As used herein the following terms refer to the structure indicated and include all of the structural isomers:

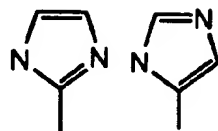
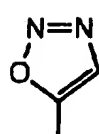
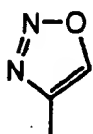
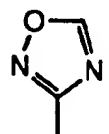
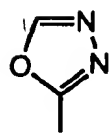
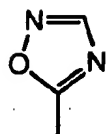
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Thiazoles

Isoxazoles

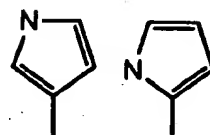
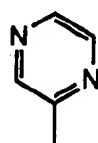
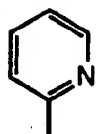
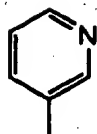
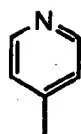
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Oxadiazoles

Imidazoles

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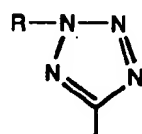
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Pyridines

Pyrazine

Pyrroles

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where R is hydrogen,
C₁-C₃ alkyl or phenyl

Tetrazole

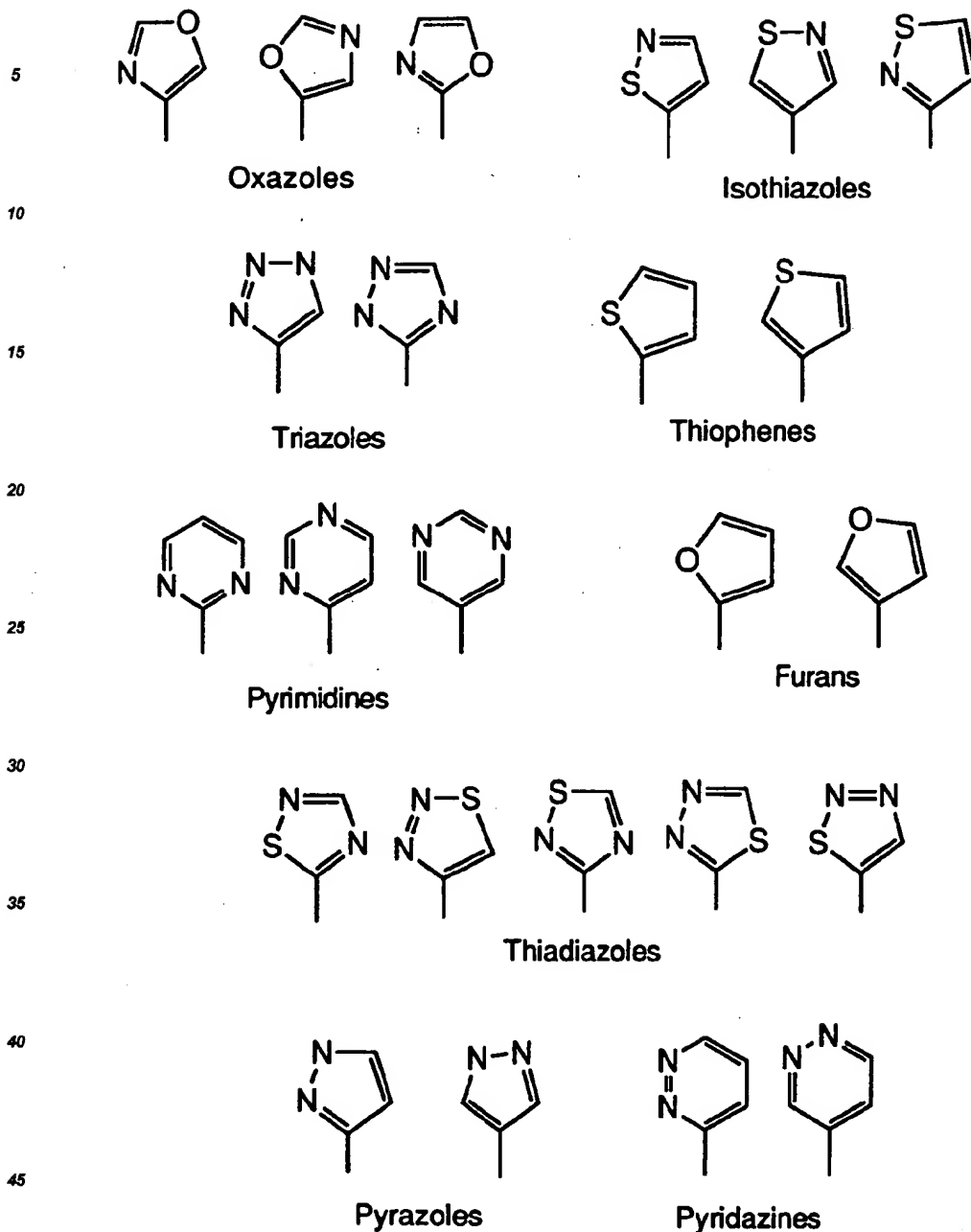
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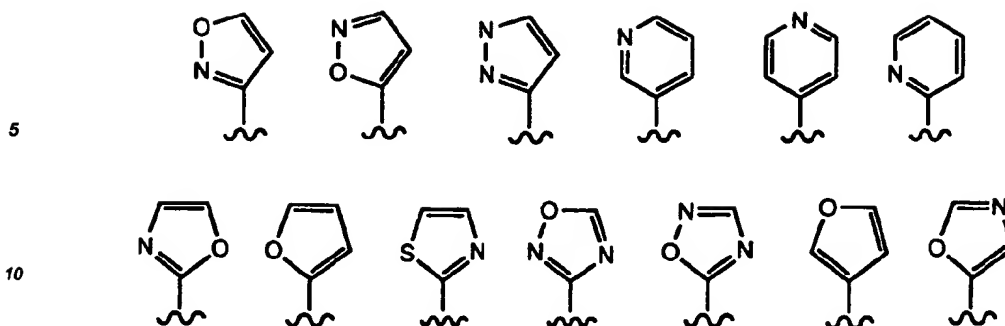
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While all of the compounds described herein are believed useful for the method of treating emesis and motion sickness provided herein, certain of such compounds are preferred for such use. Preferably R¹ and R² are both C₁-C₄ alkyl, particularly n-propyl, R³ is hydrogen, and HET is one of the following isoxazole, oxazole, pyrazole, pyridine, thiazole, furan, thiophene or oxadiazole. Other preferred aspects of the present invention are noted hereinafter.



15 The compounds of the instant invention, and the compounds employed in the method of the present invention, have at least one chiral center and therefore at least two stereoisomers can exist for each. A chiral center exists at position 4 of Formula 1. If a substituent group contains a chiral center, then additional stereoisomers can exist. Racemic mixtures as well as the substantially pure stereoisomers of Formula 1 are contemplated as within the scope of the present invention. By the term "substantially pure", it is meant that at least about 90 mole percent, more preferably at least about 95 mole percent and most preferably at least 98 mole percent of the desired stereoisomer is present compared to other possible stereoisomers. Particularly preferred stereoisomers of Formula 1 are those in which the configuration of the chiral center at position 4 is R.

20 The terms "R" and "S" are used herein as commonly used in organic chemistry to denote specific configuration of a chiral center. The term "R" refers to "right" and refers that configuration of a chiral center with a clockwise relationship of group priorities (highest to second lowest) when viewed along the bond toward the lowest priority group. The term "S" or "left" refers to that configuration of a chiral center with a counterclockwise relationship of group priorities (highest to second lowest) when viewed along the bond toward the lowest priority group. The priority of groups is based upon their atomic number (heaviest isotope first). A partial list of priorities and a discussion of stereo-chemistry is contained in the book: The Vocabulary of Organic Chemistry, Orchin, et al., John Wiley and Sons Inc., publishers, page 126, which is incorporated herein by reference.

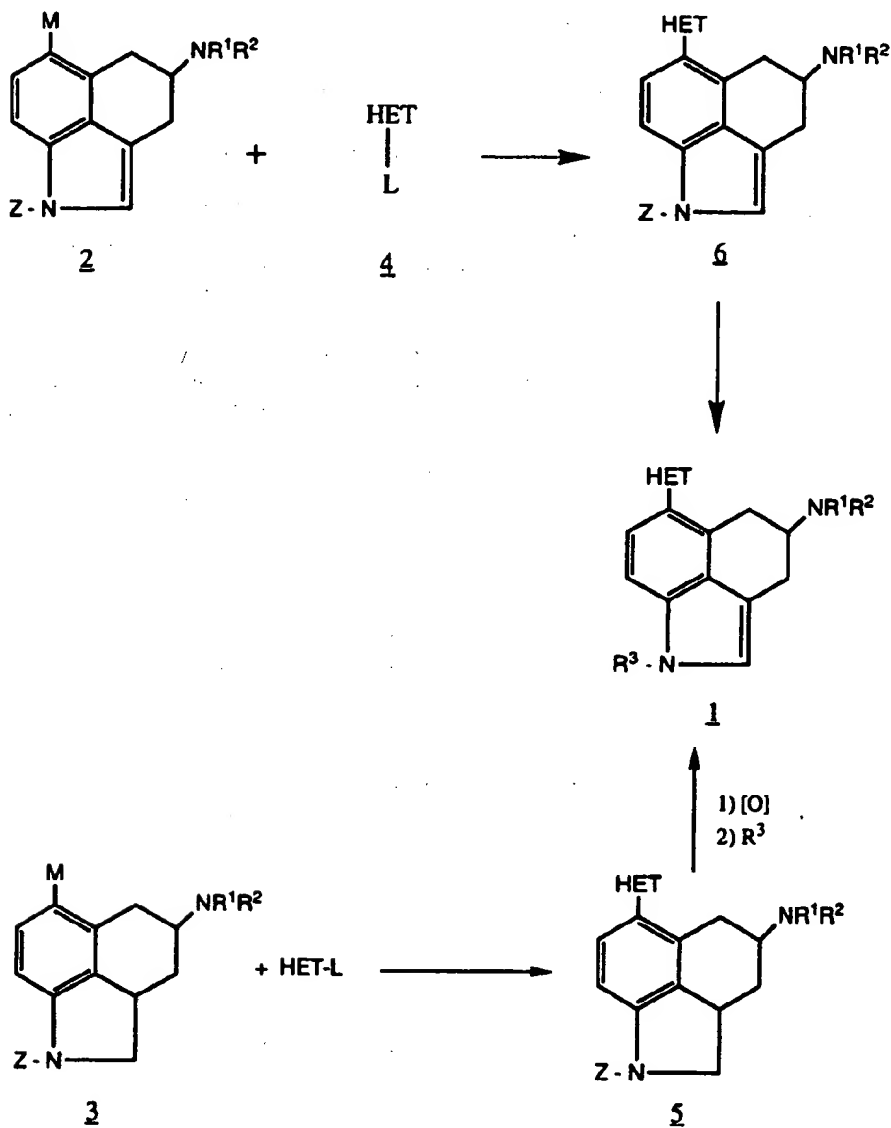
25 As set forth above, this invention includes the pharmaceutically acceptable salts of the compounds of Formula 1. Since the compounds of Formula 1 are amines, they are basic in nature and accordingly react with any number of inorganic and organic acids to form pharmaceutically acceptable salts using acids such as hydrochloric acid, nitric acid, phosphoric acid, sulfuric acid, hydrobromic acid, hydroiodic acid, phosphorous acid and others, as well as salts derived from nontoxic organic acids such as aliphatic mono and dicarboxylic acids, amino acids, phenyl-substituted alkanic acids, hydroxyalkanoic and hydroxyalkandioic acids, aromatic acids, aliphatic and aromatic sulfonic acids. Such pharmaceutically acceptable salts thus include sulfate, pyrosulfate, bisulfate, sulfite, bisulfite, nitrate, phosphate, monohydrogenphosphate, dihydrogenphosphate, metaphosphate, pyrophosphate, chloride, bromide, iodide, acetate, propionate, caprylate, acrylate, formate, tartrate, isobutyrate, caprate, heptanoate, propionate, oxalate, malonate, succinate, suberate, sebacate, fumarate, malate, mandelate, butyne-1,4-dioate, hexyne-1,6-dioate, hippurate, benzoate, chlorobenzoate, methylbenzoate, phthalate, terephthalate, benzenesulfonate, toluenesulfonate, chlorobenzenesulfonate, xylenesulfonate, phenylacetate, phenylpropionate, phenylbutyrate, citrate, lactate, β -hydroxybutyrate, glycolate, malate, naphthalene-1-sulfonate, naphthalene-2-sulfonate and mesylate.

30 Preferred compounds of Formula 1 for use in the method of treating emesis and motion sickness disclosed herein include the compounds in which R^3 is hydrogen, R^1 and R^2 are both either n-propyl or methyl and HET is 3-isoxazolyl, 5-isoxazolyl, 2-oxazolyl, 5-oxazolyl, 3-isothiazolyl, 5-isothiazolyl, 2-imidazolyl or 4-imidazolyl. These compounds include the stereoisomers at position 4, i.e., racemic mixture of 4-R and 4-S as well as the substantially pure stereoisomers of each in which the configuration at position 4 is R or S. Compounds having 4S stereochemistry are particularly preferred.

35 As depicted in Scheme I, the compounds used in the method of the present invention, as well as the compounds of the present invention, can be prepared by reacting a 4-amino-6-metallo-substituted tetrahydrobenz[cd]indole as represented by structure 2 with a heterocyclic compound represented by structure 4. In structure 2, M represents a metallo moiety such as lithium, magnesium, zinc, tin, mercury, boronic acid ($-BO_2H_2$) and the like while Z is an amino-blocking group. When the metallo moiety is multivalent, it is normally associated with other moieties such as, for example, halo for magnesium (Grignard reagent) and alkyl groups for tin (trialkyltin). The heterocycle represented by structure 4 contains a leaving group "L", such as a chloro, bromo, or trifluoromethylsulfonyl group, which can be displaced by the metalloindole. The heterocycle can be sub-

stituted as set forth hereinabove. Alternatively, a 6-metallo-1,2,2a,3,4,5-hexahydrobenz[cd]indole 3 can be similarly reacted with a heterocyclic 4 to provide the 6-heteroaryl-1,2,2a,3,4,5-hexahydrobenz[cd]indole 5 which can then be oxidized and the blocking group Z exchanged or replaced to provide the compounds of Formula 1.

Scheme 1



The reaction of the metallo-indole 2 or metalloindoline 3 and heterocycle 4 is accomplished in the presence of a palladium or nickel catalyst such as Pd[P(C₆H₅)₃]₄, PdCl₂, Pd[P(C₆H₅)₃]₂Cl₂, Ni(acac)₂, NiCl₂[P(C₆H₅)₃]₂ and the like, wherein "acac" represents acetylacetonate and "C₆H₅" represents a phenyl group. The organometallic reagent 2 or 3 is prepared by methods commonly used in the art for such preparations, for example, the lithium or magnesium reagents can be prepared by contacting the appropriate 6-chloro-, 6-bromo- or 6-iodo-substituted tetrahydro- or hexahydro- benzindole with an organolithium reagent or magnesium metal in a sol-

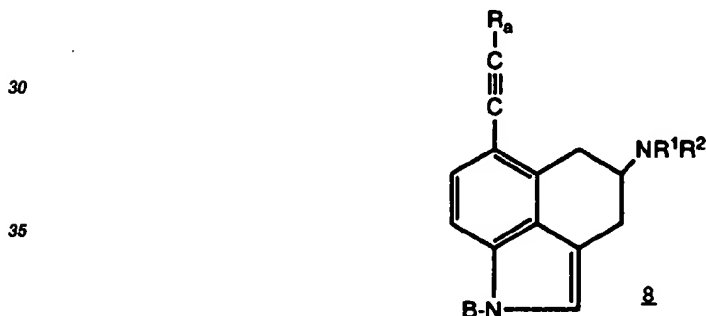
vent such as ether or tetrahydrofuran. Other organometallic derivatives can be used such as zinc, tin, mercury or boronic acid ($-\text{BO}_2\text{H}_2$). The zinc, tin and mercury reagents can be prepared by reaction of the lithiated benzindole with a zinc, tin or mercury derivative such as zinc chloride, chlorotrialkylstannane, or mercuric chloride. The boronic acid derivative can be prepared by reacting the lithium reagent with trimethylborate followed by hydrolysis of the resulting boronate ester. Mercuric acetate can be contacted directly with the hexahydrobenzindole to provide the mercurated derivative.

The 1-nitrogen of the benzindole is preferably protected with a group such as triphenylmethyl (trityl), benzyl, or, for the tetrahydrobenzindole only, triisopropylsilyl. These protecting groups are represented by Z in structures 2 and 3. The protecting group can be removed after the coupling reaction is accomplished to provide the 1-hydrobenzindole compound.

An alternative method of preparing the compounds of Formula 1 involves contacting an organometallic reagent prepared from a heterocyclic compound with a 6-bromo or 6-iodo-4-aminobenzindole. The reaction is accomplished in the presence of a catalyst such as that used in reaction Scheme I. The metal in the organometallic derivative of the heterocycle can be lithium, magnesium (Grignard reagent), zinc, tin, mercury, or a boronic acid ($-\text{BO}_2\text{H}_2$). These organometallic compounds can be prepared by standard methods, as described above for the benzindoles. Alternatively, the lithiated heterocycles can be prepared by treating a heterocycle with a strong base such as an alkyllithium or a lithium dialkylamide.

Unless otherwise indicated, in the following preparation procedures, R_a and R_a' may independently be hydrogen, $\text{C}_1\text{-C}_3$ alkyl, halogen, OH, $\text{O}(\text{C}_1\text{-C}_3 \text{ alkyl})$, $\text{S}(\text{C}_1\text{-C}_3 \text{ alkyl})$, NH_2 , CN, or phenyl. R_b may be hydrogen, $\text{C}_1\text{-C}_3$ alkyl, phenyl, or $(\text{C}_1\text{-C}_3 \text{ alkyl})\text{phenyl}$. R_c may be hydrogen or $\text{C}_1\text{-C}_3$ alkyl. R_d may be OH, $\text{O}(\text{C}_1\text{-C}_3 \text{ alkyl})$, $\text{O}(\text{phenyl})$, $\text{O}(\text{C}_1\text{-C}_3 \text{ alkylphenyl})$, halo, $\text{S}(\text{C}_1\text{-C}_3 \text{ alkyl})$, $\text{S}(\text{phenyl})$, $\text{S}(\text{C}_1\text{-C}_3 \text{ alkylphenyl})$, NH_2 , $\text{NH}(\text{C}_1\text{-C}_3 \text{ alkyl})$, $\text{N}(\text{C}_1\text{-C}_3 \text{ alkyl})_2$, $\text{OCO}(\text{C}_1\text{-C}_3 \text{ alkyl})$, $\text{OCO}(\text{phenyl})$, $\text{OCO}(\text{C}_1\text{-C}_3 \text{ alkylphenyl})$ and the like.

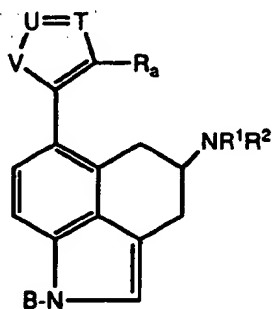
In an alternative preparation procedure, compounds of Formula 1 having a 5-membered heterocyclic ring in the 6-position can be prepared by the cycloaddition of a compound of the type represented in structure 8 wherein R^1 and R^2 are as defined above and B is an amino-protecting group or hydrogen,



with a 1,3-dipole of the type $\text{T}=\text{U}-\text{V}$ in which T, U, and V can be selected from the following list of (a) through (i).

	T	U	V
(a)	CR _a	N	CHR _a
(b)	CR _a	N	NR _b
(c)	CR _a	N	O
(d)	N	N	O
(e)	CR _a	CR _a '	NR _b
(f)	CR _a	CR _a '	O
(g)	N	CR _a '	CHR _a
(h)	N	CR _a	NR _b
(i)	N	CR _a '	O

In this list R_a and R_a' are not OH or NH₂, N represents nitrogen and O represents oxygen. This cycloaddition provides products of the structure 10, wherein R¹ and R² are as defined above and B is an amino protecting group or hydrogen.

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The 1-nitrogen of structures 8 and 10 can be protected using standard protecting groups preferably (C₂H₅)₂NC(O)-, triisopropylsilyl, or benzenesulfonyl.

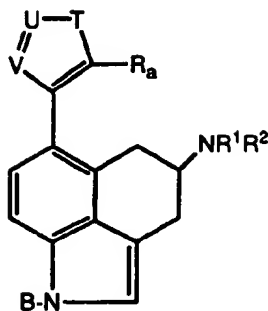
Alternatively, the 6-alkyne-substituted indole of structure 8 can be reacted with a dipole of the type ⁺T-U=V⁻ in which T, U, and V are selected from the following list for (j) and (k):

	T	U	V
(j)	CHR _a	N	N
(k)	NR _b	N	N

In this list R_a is not OH or NH₂ and N is nitrogen. This reaction provides products of structure 12.

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wherein R^1 , R^2 , R_a and B are as defined above.

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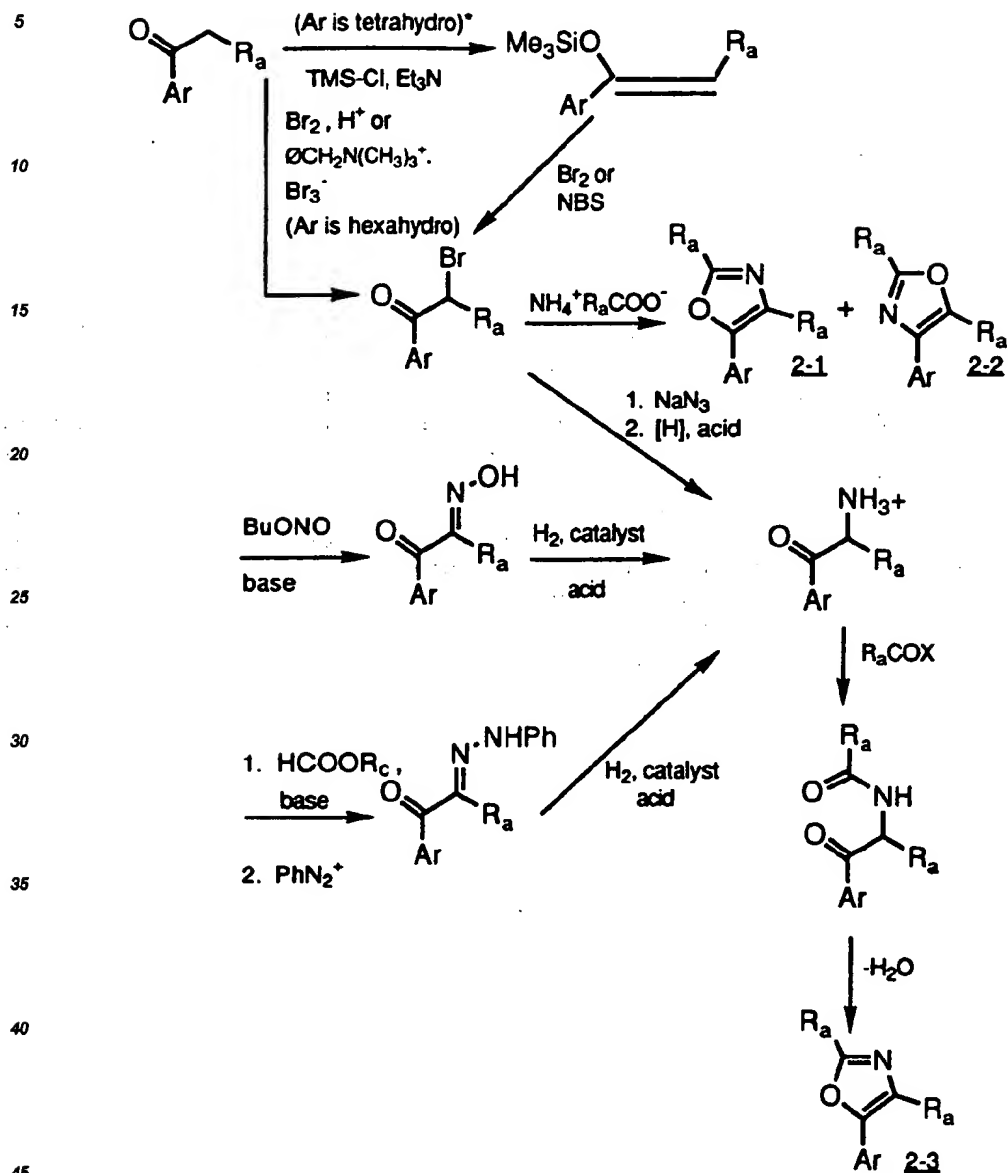
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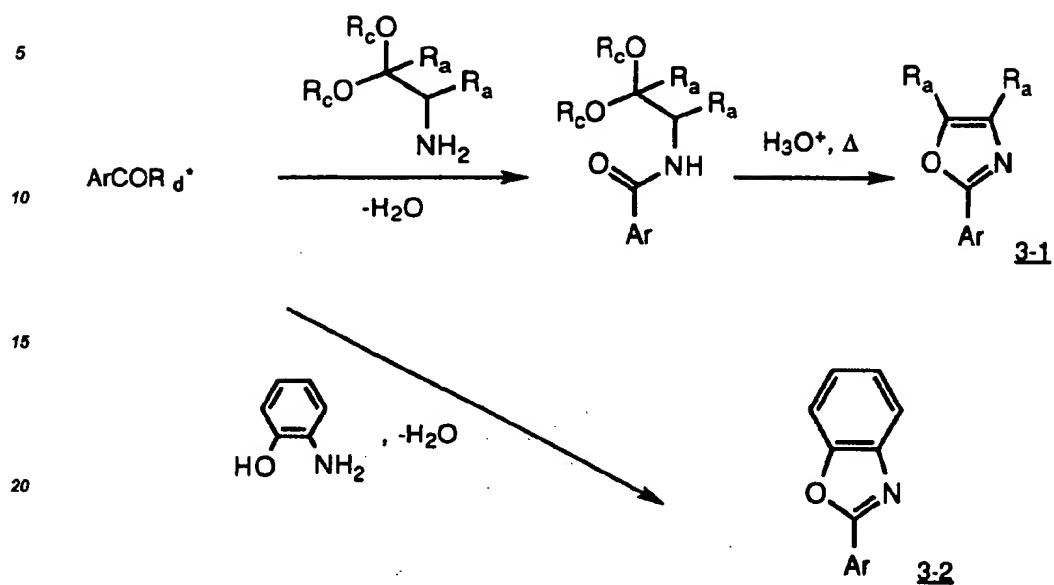
Alternative procedures for preparing certain of the compounds useful in the method of the present invention are set forth hereinbelow in Schemes 2 through 19. Scheme 19, in particular, discloses processes for preparing the novel tetrazolyl or substituted tetrazolyl compounds of the present invention. As used in these reaction Schemes, "Ar" refers to the benz[cd]indole, which can be 1,3,4,5-tetrahydro or 1,2,2a,3,4,5-hexahydro, with the indicated substituent in the 6-position, "Me" is methyl, "Et" is ethyl, "NBS" represents n-bromosuccinimide, R_a , R_b , R_c and R_d are defined above, "MsCl" represents methanesulfonyl chloride, " Δ " represents heat, " ϕ " and "Ph" each represent phenyl, "DMF" represents dimethylformamide, "TMS" represents trimethylsilyl, "[O]" represents an oxidant, Lawesson's reagent is p-methoxyphenylthionophosphine sulfide dimer, "Ac" represents acetyl, "NCS" represents N-chlorosuccinimide, "DCC" represents dicyclohexylcarbodiimide, "DMS" represents dimethyl sulfide, "Im" represents 1-imidazolyl, and "[H]" represents a reductant. As set forth hereinabove, the 1-nitrogen of the benz[cd]indole is normally protected with an amino blocking group. When Ar is tetrahydrobenz[cd]indole, the 1-blocking group is preferably triisopropylsilyl. When Ar is hexahydrobenz[cd]indole, the preferred 1-blocking group is triphenylmethyl. In the reaction schemes provided, Ar is preferably hexahydrobenz[cd]indole with the resulting 6-heteroaryl-substituted hexahydrobenz[cd]indole being oxidized to the corresponding tetrahydrobenz[cd]indole product.

Scheme 2



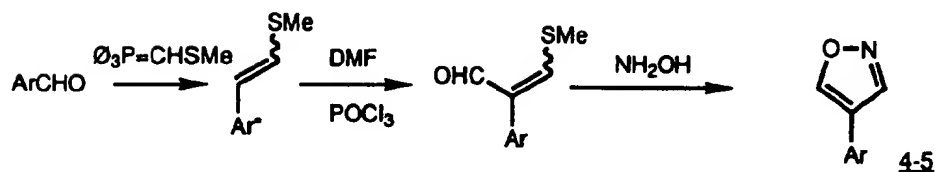
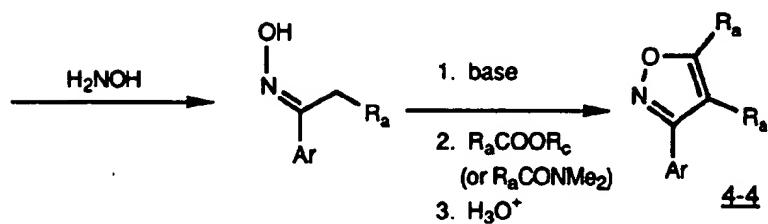
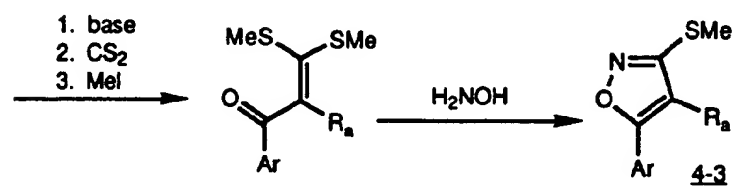
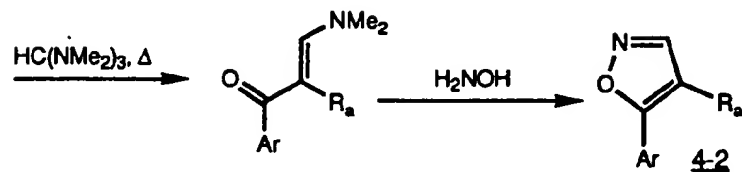
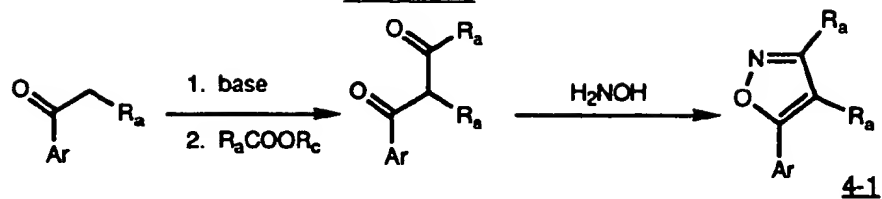
* Direct bromination can be used when Ar is tetrahydro
when the 1-nitrogen is blocked with triisopropylsilyl.

Scheme 3



* When R_d is OH the ArCOR_d substrate is preferably activated by prior contact with DCC or diimidazolylcarbonyl.

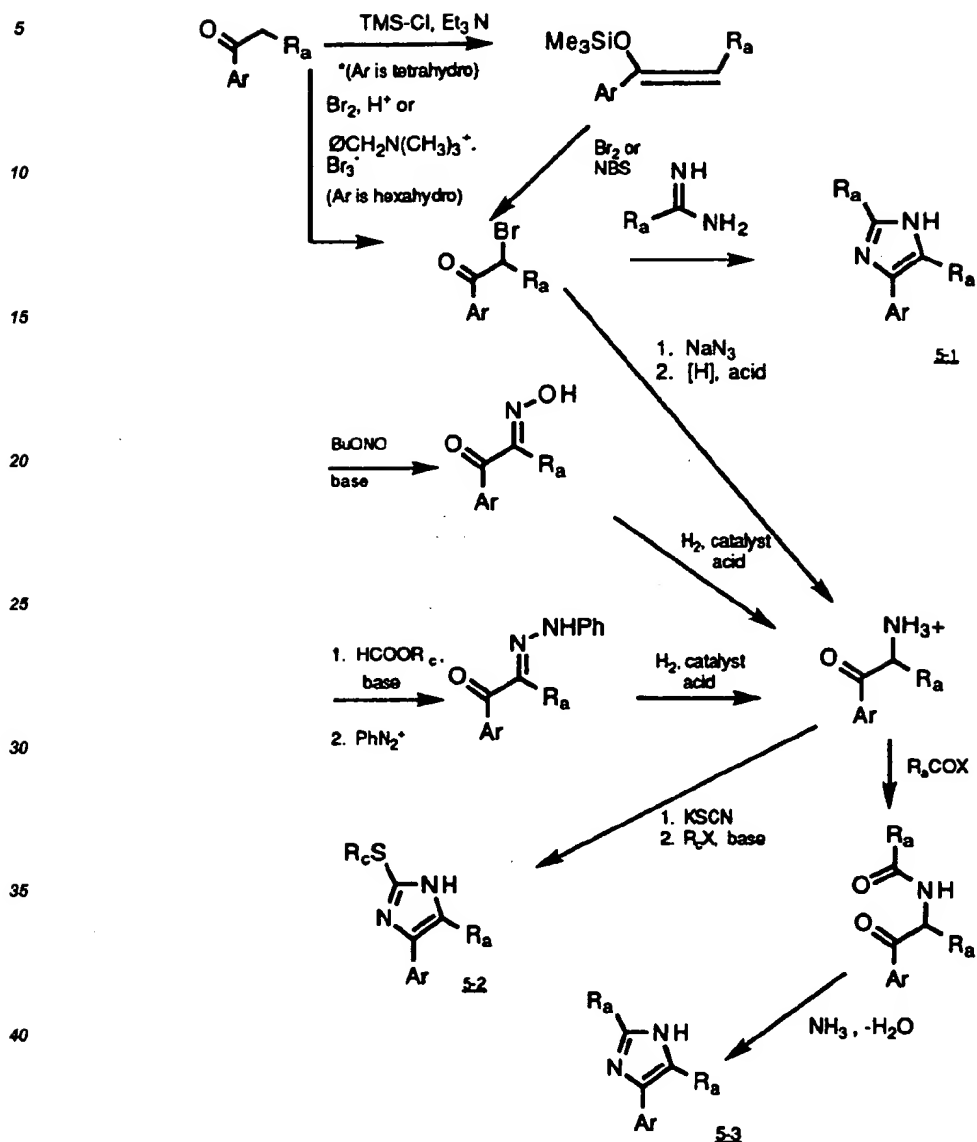
Scheme 4



* When Ar is tetrahydrobenz[cd]indole the 1-nitrogen is blocked with triisopropylsilyl.

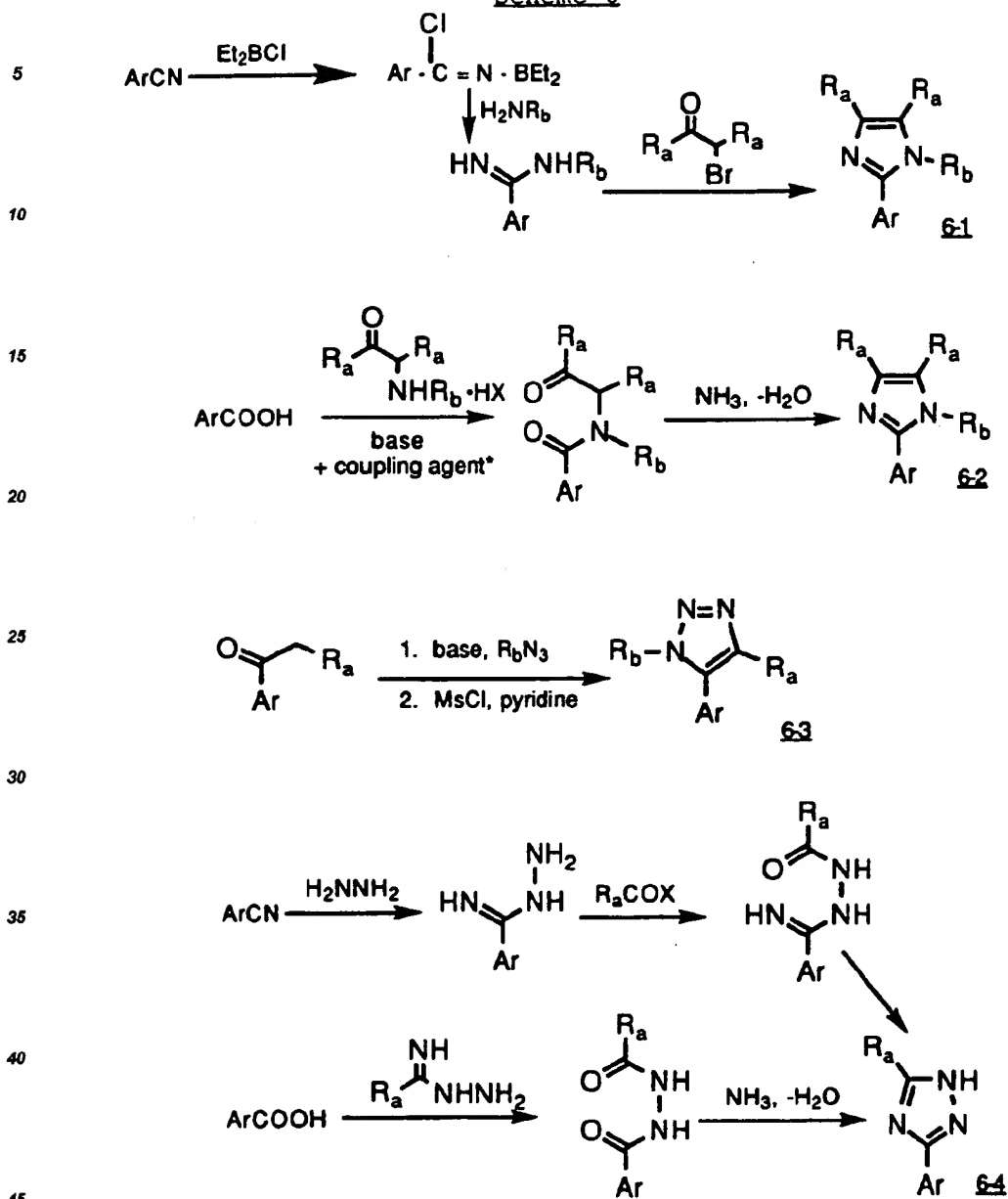
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Scheme 5



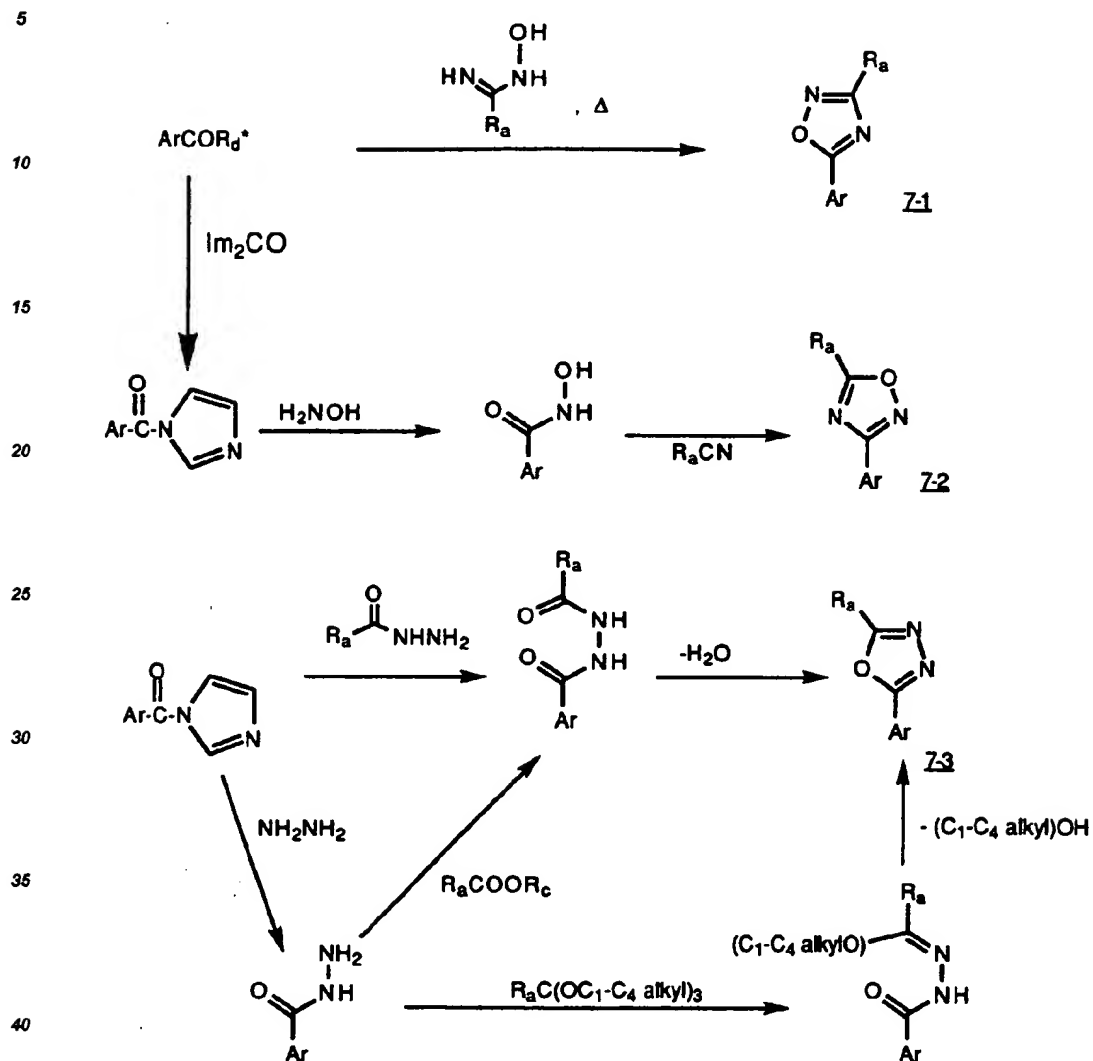
* Direct bromination can be used when Ar is tetrahydro when the 1-nitrogen is blocked with triisopropylsilyl.

Scheme 6



* For example, DCC or Im_2CO .

Scheme 7



* When R_d is OH a coupling agent, for example DCC or Im₂CO, is preferably also employed.

Scheme 8

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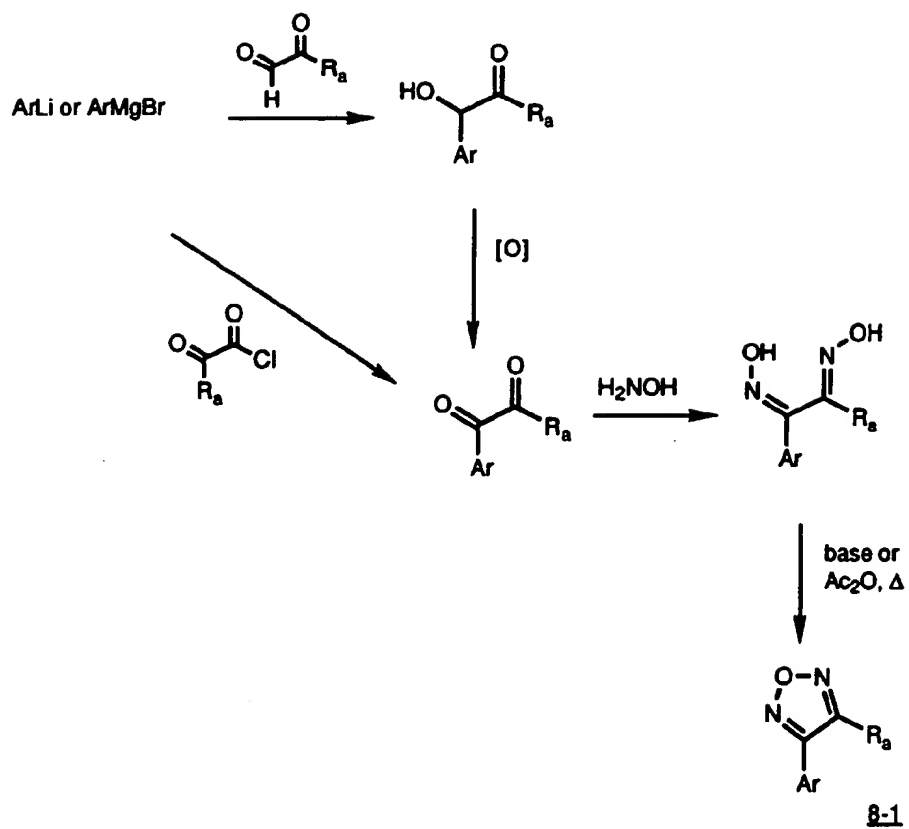
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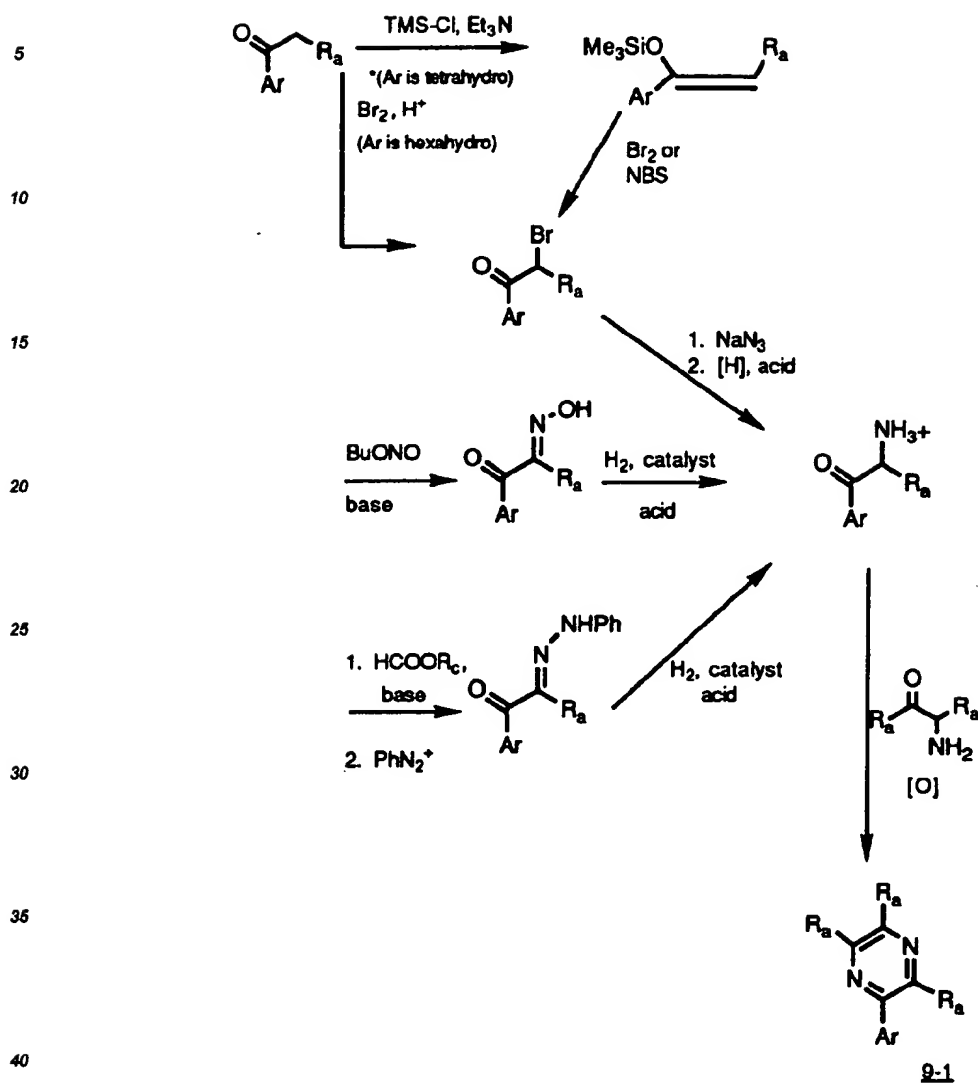
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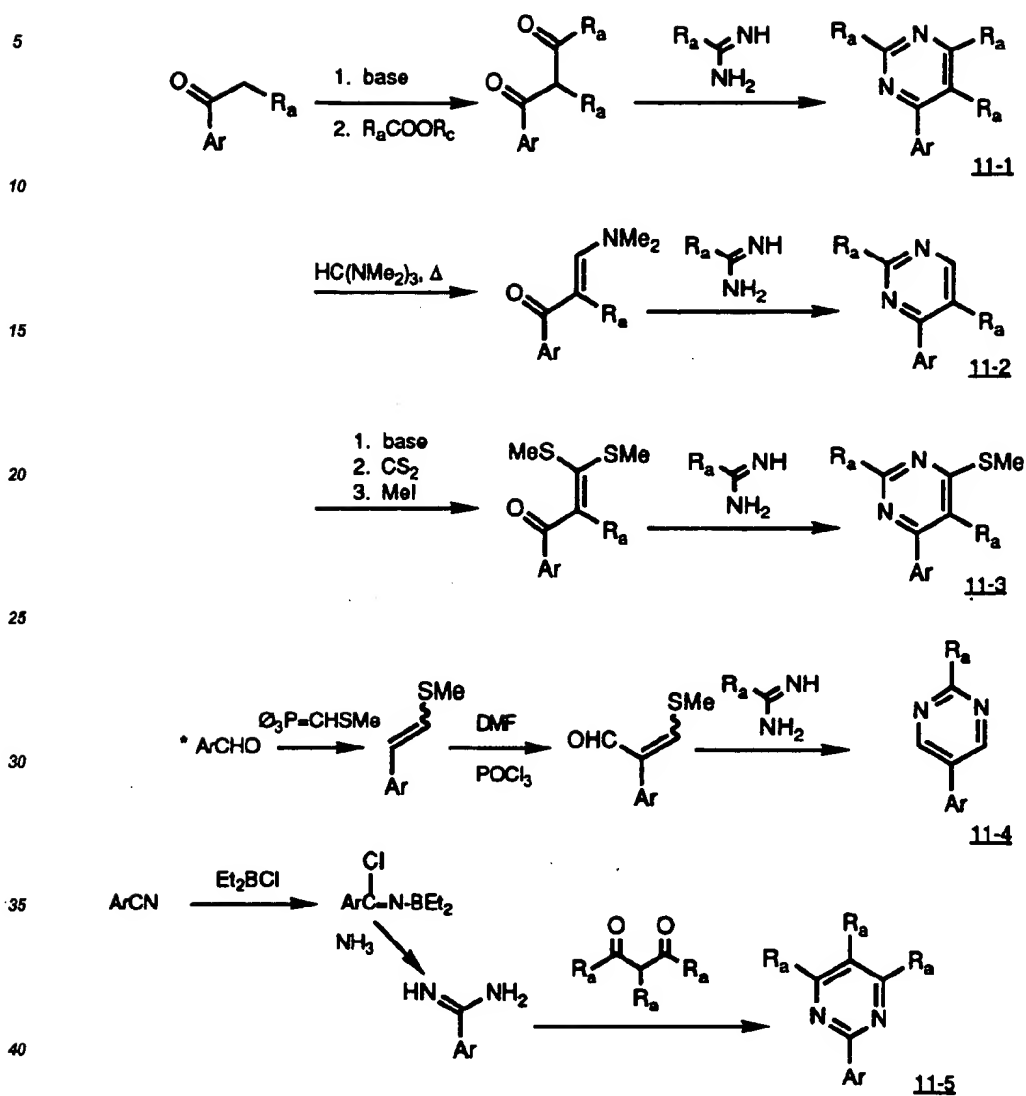
Scheme 9



*Direct bromination can be used when Ar is tetrahydro when the 1-nitrogen is blocked with triisopropylsilyl.

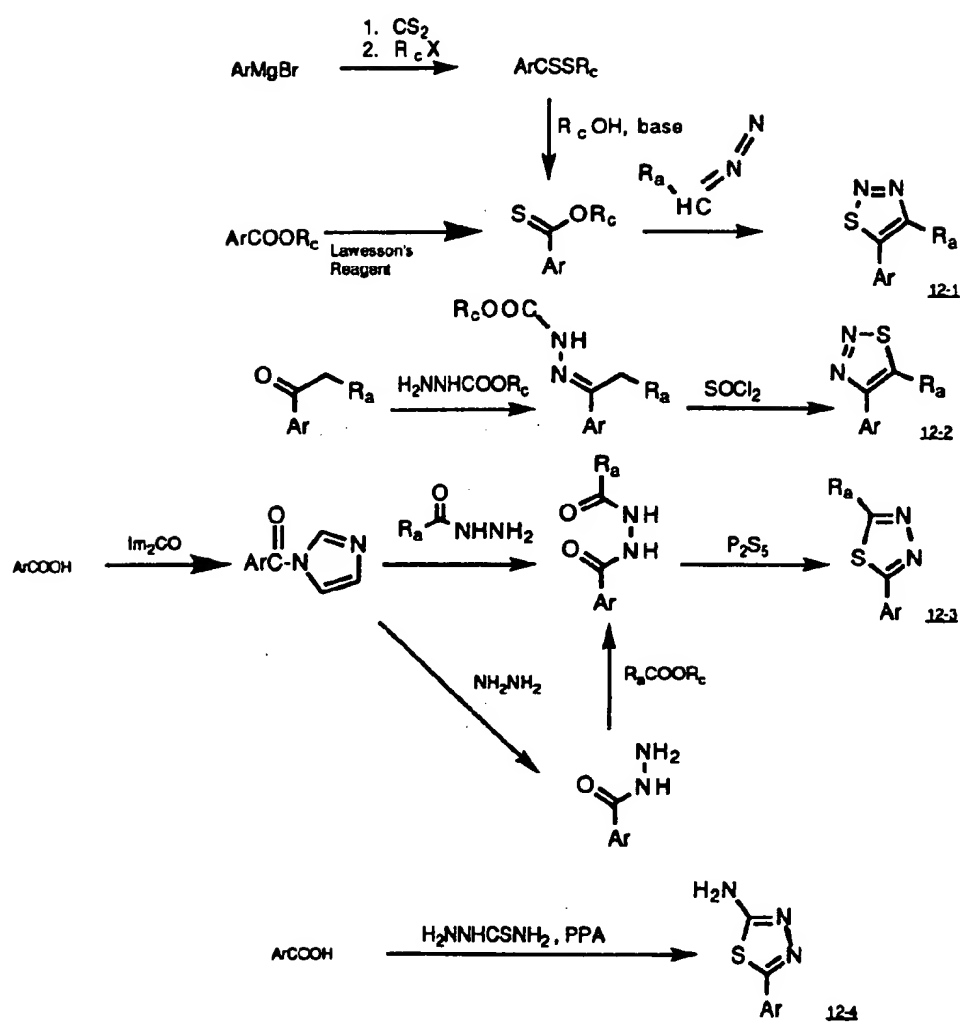


Scheme 11

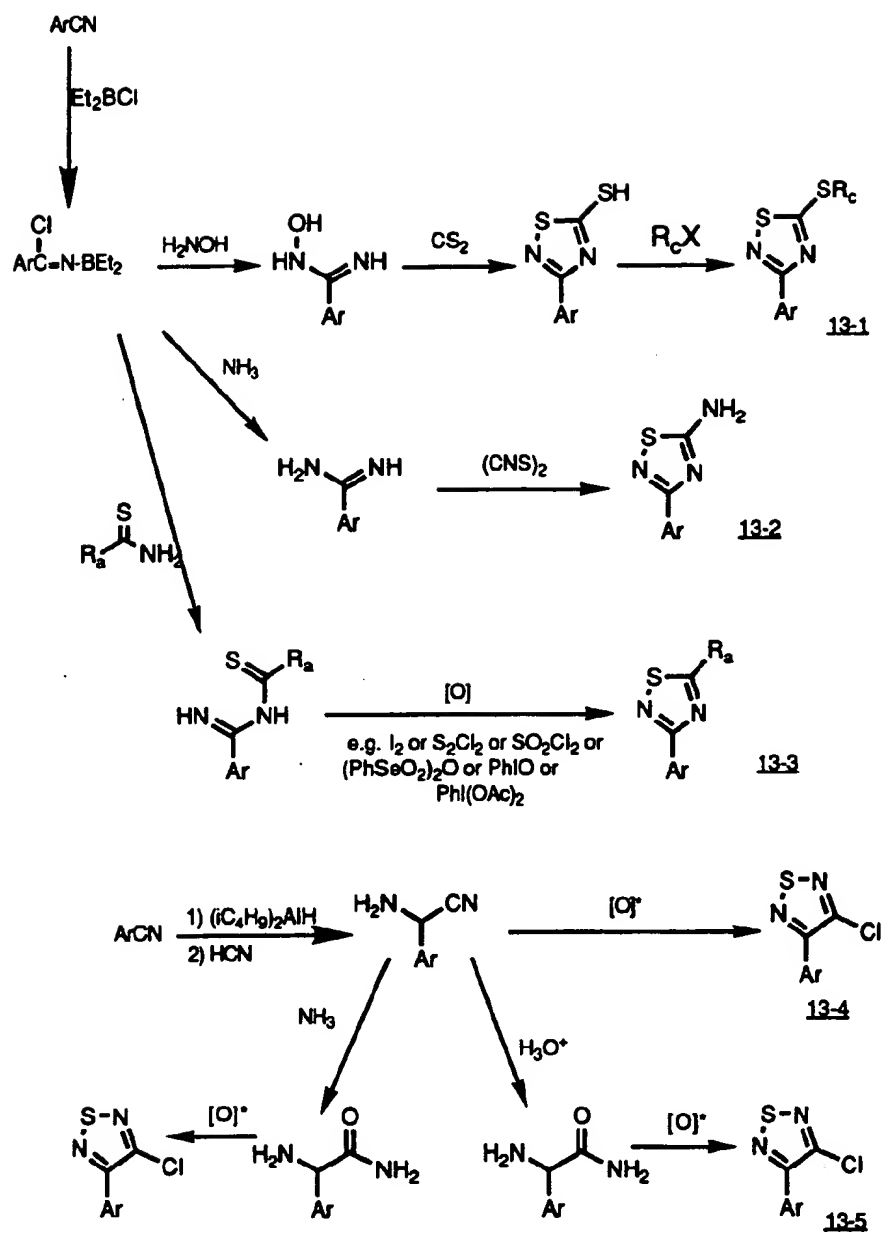


*Preferably 1-nitrogen blocked with triisopropylsilyl.

Scheme 12



Scheme 13



[O]*, e.g., SOCl₂ or SCl₂ or S₂Cl₂ or SO₂Cl₂

Scheme 14

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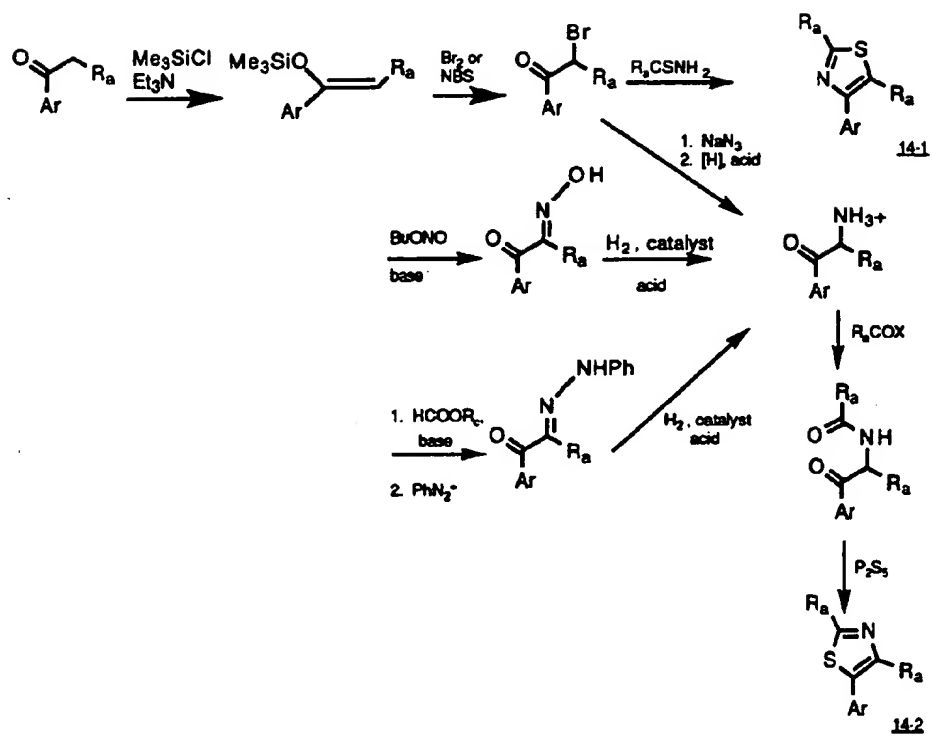
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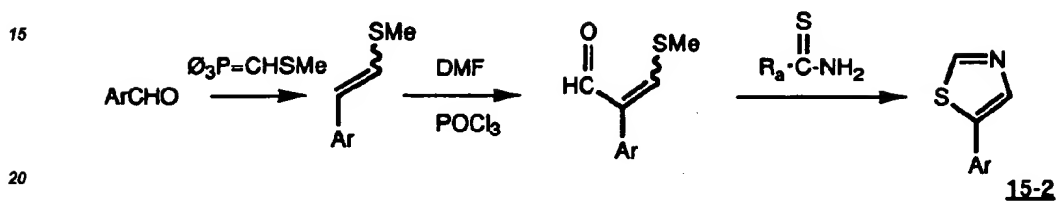
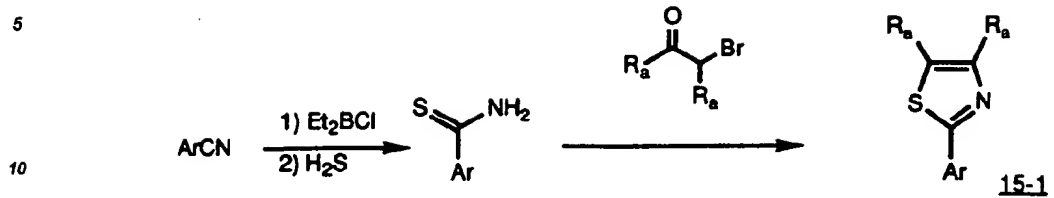
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Scheme 15



25 * 1-nitrogen preferably blocked with triisopropylsilyl.

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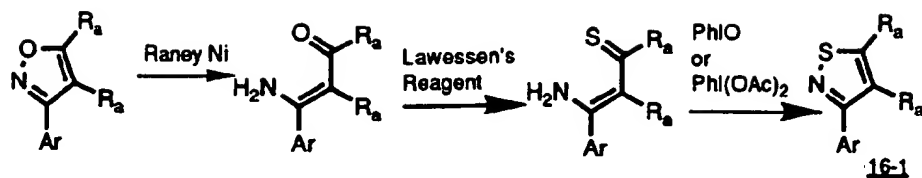
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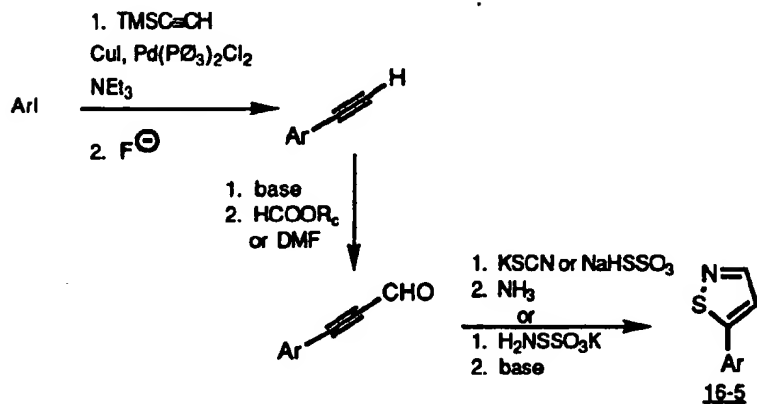
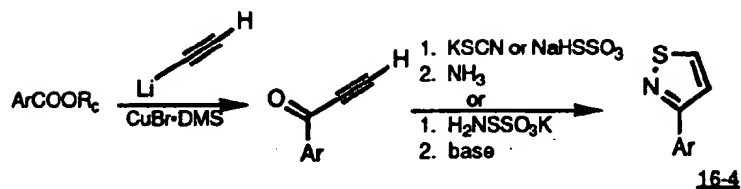
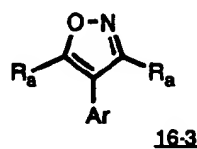
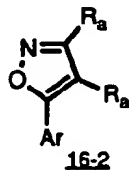
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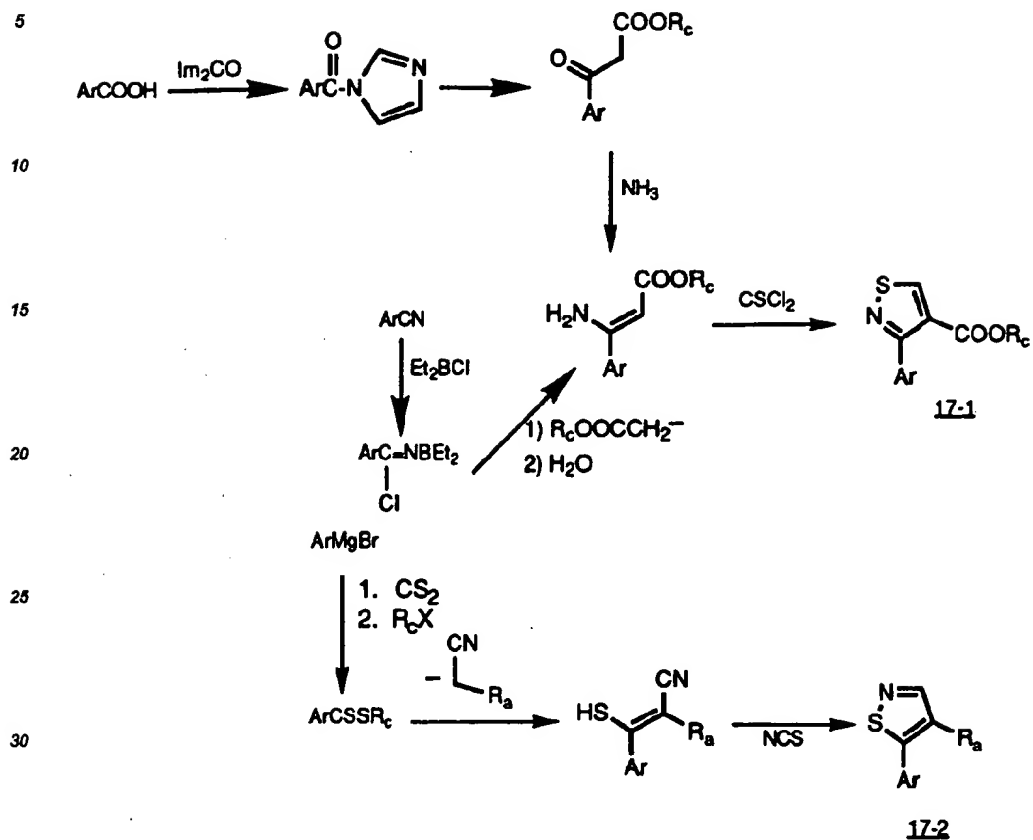
Scheme 16



15 analogous chemistry for



Scheme 17



Scheme 18

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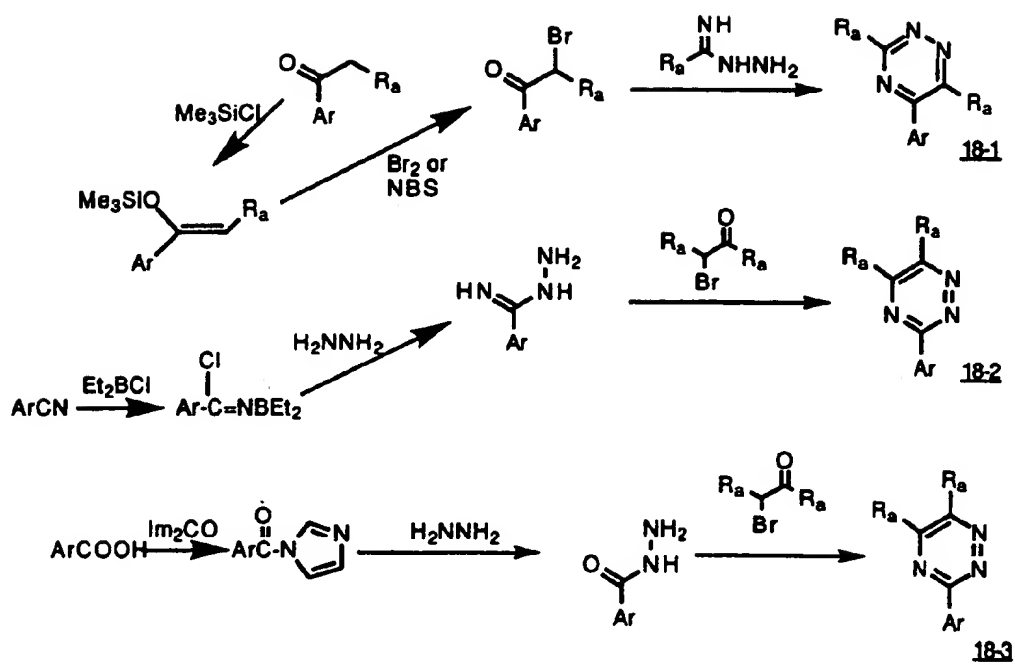
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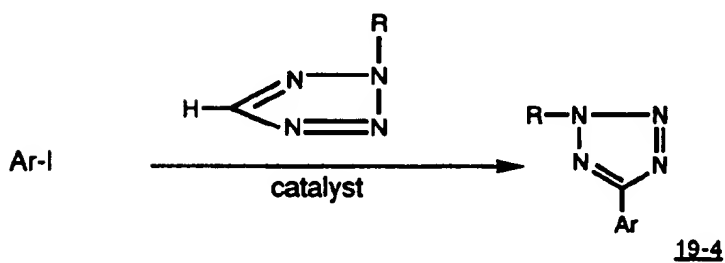
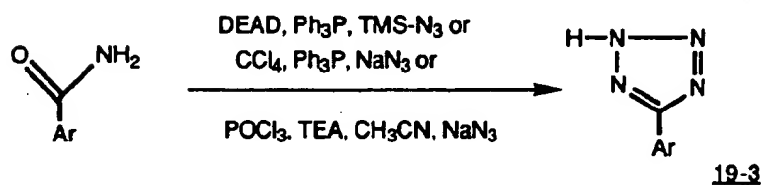
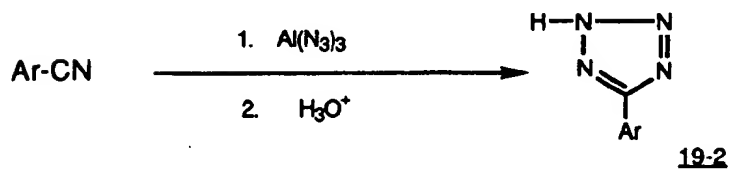
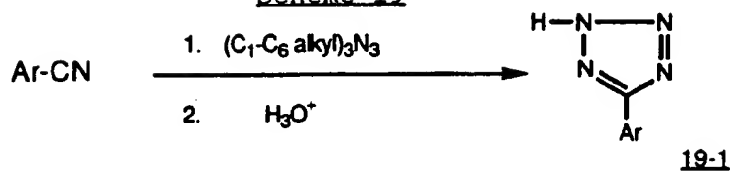
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Scheme 19



40 where R is hydrogen, C₁-C₃ alkyl or phenyl

Scheme 20 illustrates a preparation of a starting material for reaction Scheme 1.

Scheme 20

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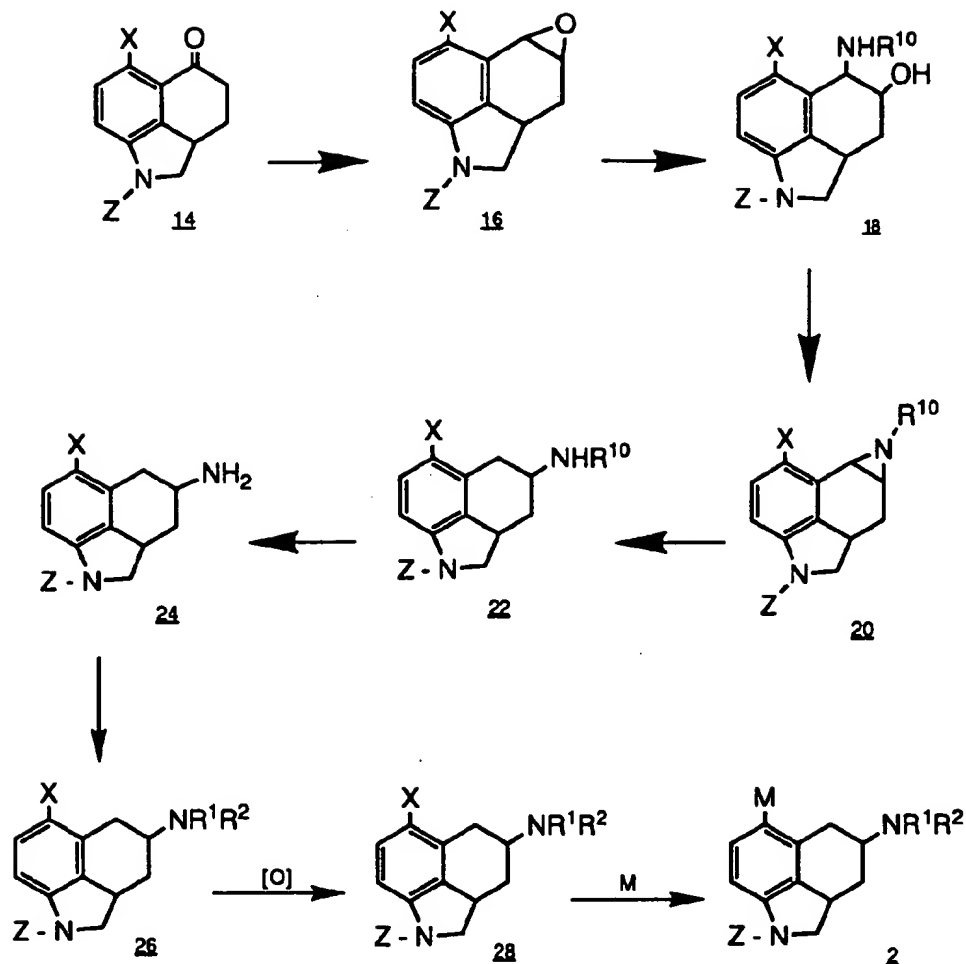
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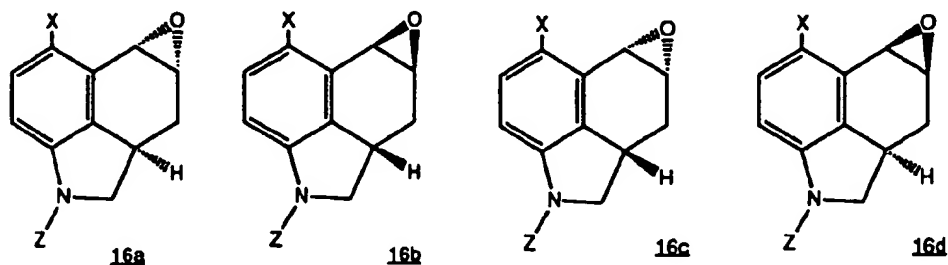


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In Scheme 20, epoxides of Formula 16 are known to the art or can be prepared from compounds known to the art using common reagents and techniques. For example, Flaugh, *et al.*, *J. Med. Chem.*, **31**, 1746 (1988); Nichols *et al.*, *Org. Prep. and Proc. Int.*, **9**, 277 (1977); and Leanna *et al.*, *Tet. Lett.*, **30**, No. 30, 3935 (1989), teach methods of preparation of various embodiments of compounds of structures 16. Those skilled in the art of organic chemistry will recognize that there are four stereoisomers of structure 16:

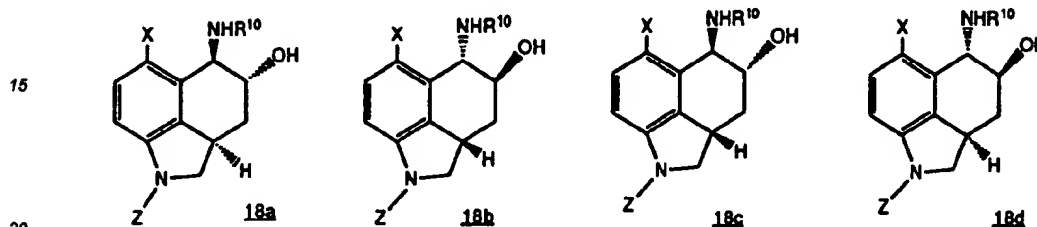
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Structures 16a and 16b are herein referred to collectively as the exo-isomers; similarly, structures 16c and 16d are the endo-isomers. Leanna et al., *supra*, teach the preparation of epoxides of structures 16 which are substantially exo or substantially endo, as desired. The preferred starting material is the compound of structure 16 wherein Z is benzoyl and X is hydrogen; the most preferred starting material is the mixture of substantially

Amino alcohols of structure 18 are formed by reacting an epoxide of structure 16 with an amine of formula $R^{10}NH_2$. Such amines are readily available. Opening of the epoxide ring proceeds substantially regiospecifically with the amino group at the 5-position and the hydroxyl group at the 4-position. The reaction is also stereospecific in the sense that stereoisomers of structure 18a-d are formed from, respectively, stereoisomers of structure 16a-d,



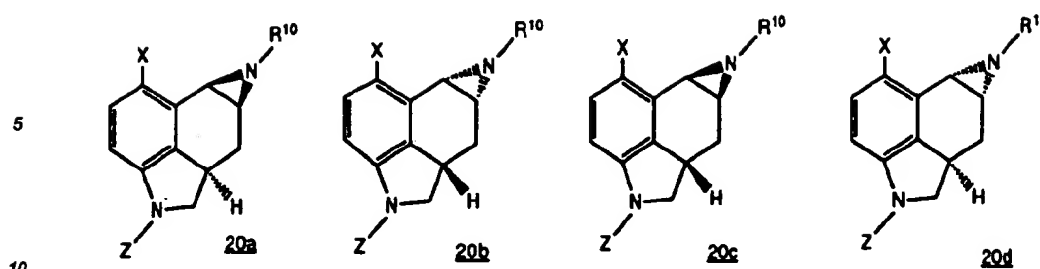
A stereoselective synthesis of the amino alcohol of structure 18, and hence of all the subsequent intermediates and products of Scheme 20, can be effected by using a substantially pure enantiomer of an amine of the formula $R^{10}NH_2$ wherein R^{10} contains at least one chiral center. The diastereomers of the resulting amino alcohol can then be separated by a number of means known in the art, for example by chromatography or crystallization. Suitable solvents for recrystallization include those such as diethyl ether, *n*-butanol, and mixtures of hexane and ethyl acetate. An alternative method of achieving a stereospecific synthesis is depicted in Scheme 20 and comprises conversion of all the diastereomers of structure 18 to corresponding diastereomers of structure 20, followed by the separation of said diastereomers of structure 20; that alternative method is discussed below. If a stereoselective synthesis is not desired, then separation of the stereoisomers of the amino alcohol of structure 18 is not required and the amine $R^{10}NH_2$ need not be optically active.

A particularly efficient stereoselective process for a highly preferred compound of structure 18, 1-benzoyl-4-hydroxy-5-(1-phenylethyl)amino-1,2,2a,3,4,5-hexahydrobenz[cd]indole, comprises the reaction of a mixture of substantially the exo-isomers of the corresponding epoxide of structure 16, or a mixture of substantially the endo-isomers of the corresponding epoxide of structure 16, with a substantially pure enantiomer of 1-phenylethylamine in the solvent *n*-butanol and the subsequent selective crystallization of one of the two isomers of the amino alcohol. The temperature of the reaction can be from about 50° to about 150°C, preferably about 80° to about 100°C.

After the reaction is complete, as determined for example by thin layer chromatography or liquid chromatography, the desired amino alcohol is crystallized at about -20° to about 40°C; the preferred temperature for the crystallization is about 0° to about 15°C. Therefore this process has the valuable attribute that the reaction and the separation of stereoisomers occur efficiently in a single step. By the proper selection of the epoxide isomers, exo or endo, and the enantiomer of 1-phenylethylamine, *R* or *S*, one can determine which of the stereoisomers of the compound of structure 18 precipitate from the reaction mixture.

A number of methods of forming aziridines such as those of structure 20 from amino alcohols such as those of Formula 18 are known to the art. Two examples are the use of diethyl azodicarboxylate and triphenylphosphine (O. Mitsunobu, *Synthesis*, January, 1981, page 1), and the use of bromine and triphenylphosphine (J. P. Freemer and P. J. Mondron, *Synthesis*, December, 1974, page 894).

A particularly efficient alternative to the above methods involves treating a compound of structure 18 with a tertiary amine in an inert solvent followed by the addition of methanesulfonyl chloride. The following stereoisomers of the aziridine of structure 20, 20a-d, arise respectively from the stereoisomers of structure 18a-d, with retention of configuration at any chiral center in the substituents Z, R^{10} or X:



Suitable tertiary amines are those of the formula $(R^{11})_3N$, where the R^{11} groups are independently C_1 - C_4 alkyl. Suitable solvents are chlorinated hydrocarbons such as methylene chloride, chloroform, carbon tetrachloride, and dichloroethane; aromatic hydrocarbons such as benzene, toluene, and the xylenes; and ethers such as tetrahydrofuran, diethyl ether, and methyl t-butyl ether. The reaction can be conducted at a temperature from about -35° to about 45°C . In the preferred embodiment, the amino alcohol is treated with triethylamine in methylene chloride at about -20° to about 0°C , then the reaction mixture is warmed to about 15° to about 35°C for the completion of the reaction. If desired, the product, an aziridine of structure 20, can be crystallized from an appropriate solvent such as acetonitrile or isopropanol after an aqueous workup. In the event that Z contains at least one chiral center in substantially a single stereoconfiguration and that the aziridine of structure 20 is prepared as a mixture of stereoisomers, said stereoisomers can be separated by methods such as chromatography and crystallization, thereby providing a stereospecific synthesis of the aziridine of structure 20 and subsequent products.

The aziridine ring can be opened to form an intermediate secondary amine of structure 22. A number of methods of opening aziridines are commonly known. It is, however, crucial that the method used for opening the aziridine to form a secondary amine of structure 22 be substantially regiospecific; the aziridine must be opened to form substantially the 4-amino compound rather than the 5-amino compound. One such method is catalytic hydrogenolysis as taught by Y. Sugl and S. Mitsui, *Bull. Chem. Soc. Jap.*, **43**, pp. 1489-1496 (1970). Catalysts which are suitable are the usual hydrogenation and hydrogenolysis catalysts, such as the noble metal catalysts; the preferred catalyst is palladium. Suitable solvents include hydrocarbons such as hexanes and heptanes; aromatic hydrocarbons such as benzene, toluene, xylenes, ethylbenzene, and t-butylbenzene; alcohols such as methanol, ethanol, and isopropanol; and mixtures of solvents such as acetic acid mixed with said alcohols. The preferred solvent for preparing the compound of structure 22, wherein Z is benzoyl, X is hydrogen, and R^{10} is 1-phenylethyl, is a mixture of tetrahydrofuran and phosphoric acid or acetic acid. The source of hydrogen can be an atmosphere of elemental hydrogen supplied at a pressure of about 1 atmosphere or higher, or the source of hydrogen can be compounds which are suitable to serve as hydrogen donors in a catalytic transfer hydrogenolysis reaction, such as formic acid, hydrazine, or cyclohexene. The preferred hydrogen source is an atmosphere of hydrogen gas supplied at about 1 to about 10 atmospheres pressure. The temperature of the reaction may be from about -20° to about 80°C ; the preferred temperature for the hydrogenolysis of the aziridine wherein Z is benzoyl, X is hydrogen, and R^{10} is 1-phenylethyl is about -20° to about 0°C .

The conversion of compounds of structure 20 to compounds of structure 22 proceeds without disturbing the stereochemical configuration of the chiral centers at the 2a- or 4-positions of the structure 22 or of the chiral centers that may be present in any of the substituents.

If desired, the compound of structure 22 can be isolated by the usual methods such as crystallization. The secondary amine of structure 22 can be converted to a primary amine of structure 24 by a number of methods known to the art of organic chemistry, or alternatively the secondary amine itself can be isolated.

However, the preferred method is to convert the secondary amine of structure 22 to the primary amine of structure 24 without isolating the secondary amine, but rather by simply continuing without interruption the hydrogenolysis reaction that produced the compound of structure 22. Therefore, the preferred solvent and catalyst are the same as those for the preparation of the secondary amine of structure 22. It may be desirable to conduct the hydrogenolysis of the secondary amine of structure 22 at a different temperature or a different pressure or different temperature and pressure than the hydrogenolysis of the aziridine of structure 20. For the hydrogenolysis of the preferred compound of structure 22 wherein Z is benzoyl, X is hydrogen, and R^{10} is 1-phenylethyl, the preferred temperature and pressure are about 50° to about 60°C and about 1 to about 20 atmospheres. Under these conditions the hydrogenolysis of compounds of structure 22 to compounds of structure 24 proceeds without disturbing the stereochemical configuration of the chiral center at the 4-position.

The isolation of the compound of structure 24 can be accomplished by the usual methods such as crys-

tallization. If desired, the compound of structure 24 can be further purified, for example by recrystallization.

Of course, as those skilled in the art will recognize, variations of Scheme 20 will be desirable or necessary for certain embodiments of the invention. For example, it may be undesirable to subject a compound in which X is halo to the catalytic hydrogenolysis steps of Scheme 19 because the undesired displacement of the halogen may compete with the desired hydrogenolysis of the carbon-nitrogen bonds. One alternative strategy is to postpone the halogenation until after the hydrogenolysis. Another alternative strategy is to use a milder means of reduction that would leave the halogen in place. A third alternative, useful in the instance when the halogen is to serve as a leaving group, is to perform the desired displacement of halogen before the hydrogenolysis step.

Compounds of Formula 1 can be prepared from the compound of structure 24, whether it exists as a mixture of stereoisomers or as a substantially pure enantiomer, using common reagents and methods well known in the art. A preferred intermediate to the compounds of the instant invention is the 6-bromo-derivative. Preferably Z is an amino blocking group such as benzoyl. A preferred method of introducing the bromo substituent at the 6-position is by reaction with bromine in glacial acetic acid, buffered with sodium acetate. Amino blocking groups can be added, if desired, to the 4-amino substituent using such methods as those disclosed by Greene, *supra*, and Barton, *supra*. Alkyl groups can be added, if desired, to the 4-amino substituent using such common methods as ammonolysis of the appropriate halide as discussed by Morrison and Boyd, Chapter 22, *Organic Chemistry*, Third Edition, Allyn and Bacon, Boston, 1973, to provide a compound of structure 26 wherein R¹ and R² are defined hereinabove. If desired, the benzoyl group can be removed from the 1-position using known methods and optionally replaced with other amino-protecting groups. Preferably the benzoyl group represented by Z is replaced with a triphenylmethyl group in structure 28 prior to the metallating step to form structure 2. The amino-protecting groups and alkyl groups can be added either before or after the bromination, as desired.

The 4-amino-6-bromotetrahydrobenz[cd]indole starting materials used to prepare the compounds of Formula 1 can be readily prepared by other processes such as depicted in Reaction Scheme 2 disclosed in United States Patent No. 4,576,959 of Flaugh, incorporated herein by reference in its entirety.

The compound of structure 26 can be oxidized to the tetrahydrobenz[cd]indole 28 using oxidizing agents such as manganese dioxide. The compound 28 where X is halo can be metallated as discussed hereinabove to provide the compound of structure 2.

The procedure of Scheme 20 using the 4,5-epoxide provides a convenient way to prepare the optically active isomers of the compounds of Formula 1. Such isomers can also be isolated by resolving racemic mixtures. This resolution can be carried out in the presence of a resolving agent, by chromatography or by repeated crystallization. Particularly useful resolving agents are d- and l-tartaric acids, d- and l-toluoyltartaric acids, and the like.

The methods of preparation described in Schemes 2-19 provide compounds in which the heteroaromatic ring may or may not be substituted. The general reactions provided below set forth methodology for incorporating, interconverting, and removing substituents on the heteroaromatic ring. Additional methods for performing these transformations are cited in *Comprehensive Organic Transformations* by Richard C. Larock, VCH Publishers, Inc., New York (1989) which is incorporated herein by reference. "HET" refers to the heteroaromatic attached to the tetrahydrobenz[cd]indole at position C-6.

1. Halogen substituent (X):

HET-OH → HET-X POX₃, PX₃, SOX₂, PPh₃X₂, or P(OR)₃X₂

HET-NH₂ → HET-X 1. HONO; 2. CuX, or KI, or HBF₄, Δ

2. O(C₁-C₃ alkyl), i.e., [OR]

HET-X → HET-OR RO-, CuI, (DMF, or DMAc, or NMP), Δ

HET-OH → HET-OR Base, RX; or CH₂N₂

3. Hydroxy substituent:

HET-NH₂ → HET-OH 1. HONO; 2. H₃O⁺, Δ

HET-OMe → HET-OH 48% HBr, Δ; or BBr₃

4. Cyano substituent:

HET-NH₂ → HET-CN 1. HONO; 2. CuCN

HET-X → HET-CN CuCN, (DMF, or DMAc, or NMP), Δ;
or CN⁻, Δ

5. S(C₁-C₃ alkyl); i.e., [SR]

HET-NH₂ → HET-SR 1. HONO; 2. RSH, base

HET-X → HET-SR RS⁻, CuI, (DMF, or DMAc, or NMP), Δ

6. Amino substituent:

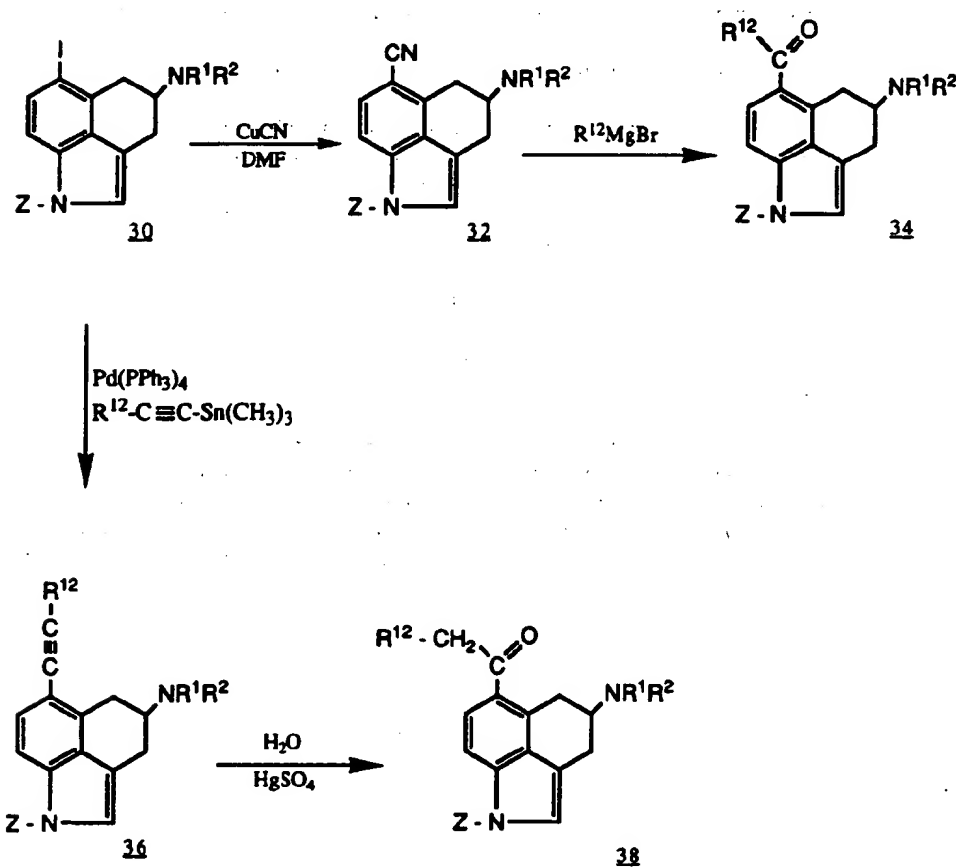
HET-NO₂ → HET-NH₂ H₂, catalyst (i.e., Pt or Pd)

7. Hydrogen substituent:

HET-X \rightarrow HET-H H_2 , catalyst; or R_3SnH , 2,2'-azobis (2-methyl)propionitrile), Δ HET-OH \rightarrow HET-H 1. 5-chloro-1-phenyltetrazole2. H_2 , catalystHET-NH₂ \rightarrow HET-H 1. HONO, 2. H_3PO_2 HET-CH₂Ph \rightarrow HET-H H_2 , catalyst (i.e., Pd) (This applies if the benzyl group is attached to a nitrogen in the heterocyclic ring.)HET-SR \rightarrow HET-H Raney Ni

6-acyl-substituted-tetrahydrobenz[cd]indoles are preferred intermediates in the preparation of certain of the compounds of Formula 1, particularly 6-isoxazole-indoles and 6-pyrazole-indoles. The 6-acyl substituted indoles can be prepared by several routes using the 6-iodo-substituted indole of structure 30 as depicted in Scheme 21 where R^1 , R^2 and Z are as defined hereinabove.

Scheme 21

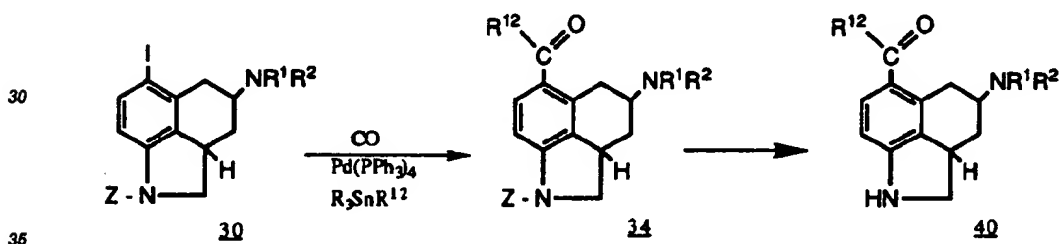


In a preferred method of preparation as depicted in Scheme 21, the nitrile 32 is contacted with an organometallic reagent such as a Grignard reagent under standard conditions to provide the 6-acyl derivative 34.

For this reaction Z is preferably triisopropysilyl. Alternatively, a 6-alkyne intermediate of structure 36 can be prepared and then hydrolyzed to provide the acyl derivative 38. This method provides a methylene group adjacent to the carbonyl group. In this method Z can be an amino protecting group such as benzoyl although the unprotected 1-nitrogen is preferred, i.e., Z is hydrogen. Compounds of structure 30 can be contacted with a palladium catalyst $\text{Pd}(\text{PPh}_3)_4$ [where Ph is phenyl] and the tin alkyne compound $\text{R}^{12}\text{-C}\equiv\text{C-Sn}(\text{CH}_3)_3$ wherein R^{12} is a $\text{C}_1\text{-C}_7$ alkyl, substituted $\text{C}_1\text{-C}_7$ alkyl, aryl ($\text{C}_1\text{-C}_3$ alkyl), substituted aryl ($\text{C}_1\text{-C}_3$ alkyl), or $\text{C}_3\text{-C}_7$ cycloalkyl. This reaction is normally conducted in a solvent such as toluene at an elevated temperature, for example at about 100°C . Typically an excess of the tin alkyne is used along with about 0.25 equivalents of the palladium compound based on compound 30. The 6-alkyne 36 is then contacted with HgSO_4 in water to provide the ketone 38. If desired the corresponding 1,2,2a,3,4,5-hexahydrobenz[cd]indole can be used in a reaction scheme similar to that of Scheme 21 followed by an oxidation step to provide structure 38. In this case the preferred blocking group for the 1-nitrogen, i.e., Z, is benzoyl or trityl.

In another preparation method depicted in Scheme 22, the 6-iodo derivative 30 can be used to prepare certain 6-acyl compounds directly. This is accomplished by contacting the 6-iodo compound with a trialkyltin alkyl complex and carbon monoxide in the presence of a palladium catalyst $\text{Pd}(\text{PPh}_3)_4$ [where Ph is phenyl] as described in the literature for arylhalides. [A. Schoenberg and R. F. Heck, *J. Org. Chem.*, 39, p. 3327 (1974); and A. Schoenberg, I. Bartoletti, and R. F. Heck, *J. Org. Chem.*, 39, p. 3318 (1974)]. Although a blocking group Z such as diethylcarbamoyl can be used for this method, the method can also be accomplished when Z is hydrogen, or the blocking group can be removed to provide compounds of structure 40 where R^1 , R^2 and R^{12} are as defined above. Alternatively the corresponding indoline can be used in the sequence of Scheme 22 followed by an oxidation step to provide structure 40.

Scheme 22



As those skilled in the art will recognize, variations of any of the reaction schemes, reagents and procedures discussed herein may be desirable or necessary for certain embodiments of the invention. Such variations are contemplated as within the scope of the present invention.

The following examples further illustrate the preparation of compounds used in the method of this invention. The examples are provided for purposes of illustration only and are not to be construed as limiting the scope of the instant invention in any way. The terms and abbreviations used in the instant examples have their normal meaning unless otherwise designated, for example, " $^\circ\text{C}$ " refers to degrees celsius; " N " refers to normal or normality; " mmol " refers to millimole; " g " refers to gram; " mL " means milliliter; " M " refers to molar; " min " refers to minutes; " hr " refers to hours; " NMR " refers to nuclear magnetic resonance; and " MS " refers to mass spectrometry.

Example 1

A. Preparation of (\pm) -1-Benzoyl-6-cyano-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

To a solution of (\pm) -1-benzoyl-6-bromo-4-(di-n-propylamino)hexahydrobenz[cd]indole (5.5 g, 12.5 mmol) in DMF (100 mL) under a N_2 atmosphere was added 3.4 g (37.5 mmol) of CuCN and 7.1 g (37.5 mmol) of CuI . The reaction mixture was then stirred at 140°C . for 8 hr. The reaction mixture was poured onto ice, diluted with water, CH_2Cl_2 added and stirred for 30 minutes. The mixture was filtered through a Celite pad and the filtrate was extracted twice with CH_2Cl_2 . The organic solution was washed twice with saturated NaCl solution. The CH_2Cl_2 solution was dried over MgSO_4 and then evaporated to provide 4 g of a solid. Chromatography of this crude product over silica gel with 1:19 $\text{MeOH}/\text{CH}_2\text{Cl}_2$ as eluent gave 3 g (62%) of product.

mp = 122-124°C.

B. Preparation of (-)(2aS,4R)-1-Benzoyl-6-cyano-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole.

To a solution of (-)(2aS,4R)-6-bromo compound (30.0 g; 0.068 mol) in 500 ml of DMF was added CuCN (18.3 g; 0.2 mol) and CuI (38.0 g; 0.2 mol). The reaction mixture was then stirred at 140°C for 6 hr. The reaction mixture was poured into 4L of water. The precipitate was collected and washed several times with water. The precipitate was suspended in dilute NH₄OH and slurried with ethyl acetate. The whole mixture was filtered through a celite pad. The ethyl acetate solution was separated and washed with brine solution. The ethyl acetate solution was dried (MgSO₄) and concentrated to dryness to provide 21.3 g of the (-)-6-nitrile.

C. Preparation of the (+)(2aR,4S)-6-cyano counterpart of Example 1B.

In a similar manner as in Example 1B above, the (+)(2aR,4S)-6-bromo compound (17.1 g, 0.039 mol) was contacted with CuCN (10.75 g; 0.12 mol) and CuI (22.8 g; 0.12 mol) in 300 ml DMF to give 11.6 g of (+)-6-cyano compound.

Example 2

Preparation of (±)-6-cyano-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole.

To a stirred solution of 4.8 g (0.0124 mol) of (±)-1-benzoyl-6-cyano-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole in 200 mL of THF cooled to -78°C under a N₂ atmosphere were added 16 mL (0.025 mol) of a 1.6 M solution of n-butyl lithium in hexane. The reaction mixture was stirred at -78°C. for 30 minutes and then allowed to warm to -20°C. To the reaction mixture was added 100 mL of 1N HCl. The mixture was extracted once with ethyl ether. The acidic solution was made alkaline with the addition of cold 5N NaOH. The basic mixture was extracted twice with CH₂Cl₂. The combined organic solution was washed with a saturated NaCl solution. The CH₂Cl₂ solution was dried over MgSO₄ and evaporated to give 4 g of an oil. Chromatography of this oil over silica gel with ethyl acetate as eluent gave 3 g (85%) of product as an oil which upon standing solidified.

Example 3

Preparation of (+)(2aS,4R)-1-trityl-6-cyano-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole.

To a solution of (+)(2aS,4R)-6-cyano-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (12.8 g, 0.045 mol) and triethylamine (4.5 g, 0.045 mol) in 400 mL of methylene chloride was added a solution of triphenylmethyl chloride (trityl chloride)(12.8 g, 0.045 mol) in 100 mL of methylene chloride dropwise at room temperature. The reaction mixture was stirred for 16 hr at room temperature. The reaction mixture was extracted with water and cold 1N HCl. The organic solution was washed with a saturated NaHCO₃ solution and with a saturated brine solution. The organic layer was dried (MgSO₄) and concentrated to dryness in vacuo to give a residue. The residue was slurried with warm hexanes, cooled and filtered to remove insolubles. The filtrate was concentrated to an oil. The oil was chromatographed (silica gel, 20% ethyl acetate in hexanes) to provide 20.6 g of the (+)-trityl nitrile.

Example 4

Preparation of (+)(2aS,4R)-6-acetyl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole.

A solution of 2.4 g (4.6 mmol) of (+)(2aS,4R)-1-trityl-6-cyano-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole in 100 mL of THF was treated with 25 mL of 2.0M methylmagnesium bromide in diethyl ether. The reaction mixture was refluxed for 16 hr. The reaction mixture was cooled and excess Grignard reagent was decomposed with addition of a saturated NH₄Cl solution. The reaction mixture was extracted with ethyl acetate. The organic solution was evaporated to an oil. The oil was dissolved in 25 mL of 5N HCl and the solution was stirred at room temperature for 30 min. The acidic solution was made alkaline with the addition of excess concentrated NH₄OH solution. The basic mixture was extracted twice with ethyl acetate. The combined organic solution was washed once with a saturated NaCl solution and dried over MgSO₄. The ethyl acetate solution was evaporated to yield 1.4 g of an oil. Chromatography of this oil over silica gel with ethyl acetate as eluent gave 1.2 g (87%) of product. Recrystallization from hexanes yielded 840 mg of the product (+) ketone.

mp = 121-122°C

[α]_D +67.40° (MeOH).

Example 5Preparation of (\pm)-6-acetyl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole.

A solution of 0.5 g (1.8 mmol) of (\pm)-6-cyano-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole in 75 mL of benzene was treated with 5 mL of 2.0M methylmagnesium bromide in diethyl ether. The reaction mixture was refluxed for 2 days. The reaction mixture was cooled and excess Grignard reagent was decomposed with addition of a saturated NH_4Cl solution. The benzene layer was separated and washed once with a saturated NaCl solution. The organic solution was evaporated to an oil. The oil was dissolved in 25 mL of 5N HCl and the solution was stirred at room temperature for 30 min. The acidic solution was made alkaline with the addition of excess concentrated NH_4OH solution. The basic mixture was extracted twice with CH_2Cl_2 . The combined organic solution was washed once with a saturated NaCl solution and dried over MgSO_4 . The CH_2Cl_2 solution was evaporated to yield 0.5 g of an oil. Chromatography of this oil over silica gel with ethyl acetate as eluent gave 0.4 g (75%) of product as an oil which upon standing solidified.

mp = 76-77° C

Example 6Preparation of (+)(2aS,4R)-6-(3-pyrazoyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole-2 HCl.

A solution of (+)(2aS,4R)-1-triphenylmethyl-6-acetyl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (1.67 g, 3 mmol) and 3 mL of tris(dimethylamino)methane in 50 mL of toluene was refluxed for 5 hr. The reaction was concentrated in vacuo and the residue was dissolved in 100 mL of CH_3OH . To the CH_3OH solution was added 2 mL of 85% hydrazine and the reaction mixture was stirred at room temperature for 16 hours. To the reaction mixture was added 50 mL of 1N HCl and stirred for an additional 1 hr. The solution was concentrated in vacuo to remove CH_3OH and the acidic solution was extracted with ethyl acetate. The acidic solution was separated and made alkaline with addition of excess concentrated NH_4OH . The basic mixture was extracted with ethyl acetate. The ethyl acetate solution was washed with a brine solution, dried (MgSO_4) and concentrated in vacuo provide 900 mg of an oil. The crude product was chromatographed through silica gel (flash column, ethyl acetate) to yield 700 mg of pyrazole compound. The oil was dissolved in 50 mL of CH_3OH and 2 equivalents of 0.1 N HCL was added to the solution. The solution was concentrated in vacuo and the residue was crystallized from ethanol/diethyl ether.

Yield - 400 mg

mp = 260 d

MS m/e 324(FD)

$[\alpha]_D = +19.84^\circ$ (MeOH).

Analysis calculated for $\text{C}_{20}\text{H}_{28}\text{N}_4 \cdot 2\text{HCl}$			
Theory:	C, 60.45;	H, 7.61;	N, 14.10;
Found:	C, 60.21;	H, 7.60;	N, 14.26.

Theory:	C, 60.45;	H, 7.61;	N, 14.10;
Found:	C, 60.21;	H, 7.60;	N, 14.26.

Example 7Preparation of (\pm)-6-(5-isoxazolyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole-2 HCl.

To a solution of (\pm)-6-acetyl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (2.3 g, 7.7 mmol) and triethylamine (1.1 mL, 8 mmol) in 90 mL CH_2Cl_2 under N_2 was added dropwise a solution of 2,2,2-trichloroethyl chloroformate. The reaction mixture was stirred at room temperature for 1 hr. The CH_2Cl_2 solution was extracted with water and 1N HCl . The organic solution was washed with a saturated NaHCO_3 solution and with a brine solution. The CH_2Cl_2 solution was dried (MgSO_4) and concentrated in vacuo to give 3.3 g of the 1-carbamylindoline.

A solution of this 1-carbamylindoline (3.3 g, 7.7 mmol) and tris(dimethylamino)methane (5 mL) in 70 mL of toluene was stirred at reflux for 16 hr. The reaction mixture was concentrated to dryness in vacuo. The residue was dissolved in 50 mL of acetic acid and hydroxylamine hydrochloride (2.5 g, 36 mmol) was added. The reaction mixture was stirred at room temperature for 16 hr and then concentrated in vacuo to dryness. The residue was suspended in water and excess concentrated NH_4OH was added to the mixture. The basic mixture was extracted with CH_2Cl_2 . The organic solution was washed with a brine solution, dried (MgSO_4) and concentrated in vacuo to give 3.1 g of an oil. The crude product was chromatographed (flash column, silica gel,

20% hexanes in ethyl acetate) to yield 2.0 g of (\pm)-1-carbamyl-6-isoxazolyindoline.

This isoxazole carbamate was dissolved in 20 mL of acetic acid and 1 g of zinc dust was added all at once. The reaction mixture was stirred at room temperature for 4 hr. The reaction mixture was filtered through a celite pad and the filtrate was concentrated to dryness in vacuo. The residue was suspended in a saturated NaHCO_3 solution and extracted with CH_2Cl_2 . The organic solution was washed with a brine solution, dried (MgSO_4) and concentrated to an oil. The crude material was chromatographed (flash column, silica gel, ethyl acetate) to give 500 mg of isoxazole indoline. The product was dissolved in 50 mL of CH_3OH and 2 equivalents of 0.1N HCl were added. The solution was concentrated to dryness and the residue was crystallized from ethanol/diethyl ether to give 85 mg of 5-isoxazole-6-substituted product as the dihydrochloride.

mp = 226°C d

MS m/e 325(FD)

Analysis calculated for $\text{C}_{20}\text{H}_{27}\text{N}_3\text{O}\cdot 2\text{HCl}$

Theory:	C, 60.30;	H, 7.34;	N, 10.55;
Found:	C, 58.83;	H, 7.18;	N, 10.01.

Example 8

Preparation of (+)(2aS,4R)-6-(3-isoxazolyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole-2 HCl.

A solution of (+)(2aS,4R)-1-triphenylmethyl-6-acetyl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (3.33 g, 6 mmol), 5 g hydroxylamine hydrochloride, 20 mL pyridine and 30 mL of ethanol was refluxed for 16 hr. The reaction mixture was concentrated to dryness in vacuo and the residue was dissolved in 5N HCl. The acidic mixture was extracted with ethyl acetate. The acidic solution was made alkaline with excess NH_4OH solution and extracted with ethyl acetate. The ethyl acetate solution was washed with a brine solution, dried (MgSO_4) and concentrated in vacuo to give 1.5 g of crude product which was chromatographed (flash column, silica gel, ethyl acetate) to give 1.2 g of oxime.
mp = 129-130°C.

To a solution of this oxime (1.2 g, 3.8 mmol) in 100 mL of THF cooled to -5°C under a N_2 atmosphere was added 7.5 mL n-butyllithium (1.6 M in hexanes) dropwise with stirring. The reaction mixture was stirred with continued cooling for 1 hr. To the reaction mixture was added 2 mL (26 mmol) of DMF all at once and then stirred for 1 hr at room temperature. The reaction mixture was poured into 50 mL of 1N H_2SO_4 and the acidic solution was warmed on a steam bath for 1 hr. The acidic solution was cooled, extracted with diethyl ether, and then made alkaline with excess 5N NaOH. The basic mixture was extracted with ethyl acetate. The organic layer was washed with a brine solution, dried (MgSO_4) and concentrated in vacuo to give 1 g of an oil. The oil was chromatographed (flash column, silica gel, ethyl acetate) to yield 500 mg of product as an oil. The oil was dissolved in 50 mL of CH_3OH and 2 equivalents of 0.1N HCL were added. The solution was concentrated to dryness in vacuo and the residue was crystallized from ethanol/diethyl ether. Crystallization gave 300 mg of the dihydrochloride of the 6-(3-isoxazolyl) product.

mp = 215°C d

MS m/e 325(FD)

$[\alpha]_D^{25} +26.4^\circ$ (MeOH).

Example 9

Preparation of (\pm)-1-benzoyl-6-[4-(2-aminothiazolyl)]-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole.

To a solution of (\pm)-6-acetyl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (205 mg, 0.7 mmol) and triethylamine (81 mg, 0.8 mmol) in 20 mL of CH_2Cl_2 was added a solution of benzoyl chloride (112 mg, 0.8 mmol) in 20 mL of CH_2Cl_2 . The reaction mixture was stirred at room temperature for 2 hr. The reaction mixture was successively washed with water, a saturated NaHCO_3 solution, a brine solution and dried (MgSO_4). The organic layer was concentrated to dryness in vacuo to give 200 mg of the 1-benzoyl derivative.

A solution of this N-benzoyl compound (200 mg, 0.5 mmol) in 20 mL of acetic acid was saturated with HBr(gas). To the solution was added dropwise a solution of bromine (0.2 mL) in 5 mL of acetic acid. The reaction was stirred at room temperature for 30 min and then concentrated to dryness in vacuo. The residue was dis-

solved in 30 mL of ethanol then 500 mg of thiourea were added and the mixture refluxed for 16 hr. The reaction was concentrated to dryness in vacuo and the residue dissolved in water. The solution was made alkaline with the addition of concentrated NH_4OH . The basic mixture was extracted with CH_2Cl_2 . The organic solution was washed with a brine solution, dried (MgSO_4) and evaporated to dryness to give 200 mg of an oil. The oil was chromatographed (flash column, silica gel, ethyl acetate) to provide 140 mg of the named 6-aminothiazolyl compound.

MS m/e 460(FD)

Example 10

Preparation of (+)(2aS,4R)-6-(5-isoxazolyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole-2 HCl

To a solution of (+)(2aS,4R)-6-acetyl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (1.7 g, 5.7 mmol) and triethylamine (0.8 ml, 6 mmol) in 90 ml CH_2Cl_2 was added dropwise a solution of 2,2,2-trichloroethylchloroformate (1.3 g, 6 mmol) in 10 ml CH_2Cl_2 . The reaction mixture was stirred at room temperature for one hour and then extracted with water and 1N HCl. The organic solution was washed with a saturated NaHCO_3 solution, a saturated brine solution, dried over MgSO_4 and then concentrated to dryness in vacuo to give 2.5 g of the 1-carbamylindoline.

A solution of the 1-carbamylindoline (2.5 g, 5.7 mmol) and tris(dimethylamino)methane (5 ml) in 100 ml of toluene was stirred at reflux for 16 hours. After 16 hours the reaction mixture was concentrated to dryness in vacuo. The resulting residue was dissolved in 50 ml of acetic acid and 1.5 g (22 mmol) of a hydroxylamine hydrochloride solution were added. The resulting reaction mixture was stirred at room temperature for 16 hours and then concentrated to dryness in vacuo. The resulting residue was suspended in water and an excess of a concentrated NH_4OH solution was added to the mixture. The basic mixture was then extracted with CH_2Cl_2 and the resulting organic extract was washed with a saturated brine solution, dried over MgSO_4 and then concentrated in vacuo to give 2.1 g of an oil. This oil was chromatographed (flash column, silica gel, EtOAc) to yield 1.9 g of (+)(2aS,4R)-6-(5-isoxazolyl)indoline. The isoxazolylindoline was dissolved in 30 ml of acetic acid and 1.5 g of zinc dust were added all at once. The resulting reaction mixture was stirred at room temperature for four hours and then filtered through a celite pad. The filtrate thus obtained was then concentrated to dryness in vacuo. The resulting residue was suspended in a saturated NaHCO_3 solution, which was then extracted with CH_2Cl_2 . The organic extract was then washed with a saturated brine solution, dried over MgSO_4 and concentrated in vacuo to an oil. This oil was chromatographed (flash column, silica gel, EtOAc) to give 400 mg of isoxazolylindoline. Such compound was dissolved in 50 ml of methanol and two equivalents of 0.1N HCl were added. The resulting solution was concentrated to dryness in vacuo and the resulting residue was then crystallized from ethanol/diethyl ether to give 170 mg of title compound.

mp = 235°C d

MS m/e 325(FD)

$[\alpha]_D + 27.29^\circ$ (MeOH)

Analysis calculated for $\text{C}_{20}\text{H}_{27}\text{N}_3\text{O} \cdot 2\text{HCl}$

Theory:	C, 60.30;	H, 7.34;	N, 10.55;
Found:	C, 60.53;	H, 7.54;	N, 10.26.

Example 11

Preparation of (-)(2aR,4S)-6-(5-isoxazolyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole-2 HCl

The title compound was prepared substantially in accordance with the method described in Example 10, above, utilizing 2.5 g (8.3 mmol) of (-)(2aR,4S)-6-acetyl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (prepared substantially in accordance with the method described in Example 4) and 1.5 g (22 mmol) of a hydroxylamine hydrochloride solution. Such reaction sequence provided 500 mg of title compound.

m.p. 235°C d

MS m/e 325(FD)

$[\alpha]_D - 29.18^\circ$ (MeOH)

Analysis calculated for C ₂₀ H ₂₇ N ₃ O·2HCl			
Theory:	C, 60.30;	H, 7.34;	N, 10.55;
Found:	C, 60.11;	H, 7.41;	N, 10.43.

Example 12

Preparation of (2aR,4S)-6-(3-phenyl-1,2,4-oxadiazol-5-yl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

A sodium ethoxide solution was prepared by dissolving 49 mg (2.1 mmol) of sodium in 35 ml of ethanol. Phenylhydroxamidine (1.73 g, 12.71 mmol) and 6-ethoxycarbonyl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (890 mg, 2.1 mmol) were added to the ethoxide solution and the resulting solution was heated to reflux and stirred at that temperature for 6.25 hours and then stirred overnight at room temperature. The next morning additional sodium ethoxide solution (50 mg of sodium in 10 ml of ethanol) was added and the reaction mixture was again stirred at reflux overnight. The next morning water was added to the reaction mixture and the resulting solution was then extracted with ethyl acetate. The organic extract was washed sequentially with water and a saturated brine solution, dried over sodium sulfate and then concentrated *in vacuo* to provide 2.33 g of a brown oil. This oil was purified by flash chromatography (2.5% isopropanol in chloroform plus 0.5% ammonium hydroxide) to provide 260 mg of title product as a light yellow solid. Such product was purified by recrystallization from hexane.

Analysis calculated for C ₂₅ H ₃₀ N ₄ O:			
Theory:	C, 74.59;	H, 7.51;	N, 13.92;
Found:	C, 74.59;	H, 7.52;	N, 13.90.

Example 13

Preparation of (-)-(2aR,4S)-6-(2-furyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

To a sealable tube with threads containing 13 ml of dry tetrahydrofuran were added 1.2 g (2.46 mmol) of (+)-(2aR,4S)-1-benzoyl-6-iodo-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole, 968 mg (2.71 mmol) of 2-(tributylstannyl)furan and 200 mg of bis(triphenylphosphine)palladium(II) chloride. The resulting mixture was then deaerated with argon for 15 minutes. After deaeration, the tube was sealed with a teflon cap and the contents thereof were heated to 100°C for 24 hours. After 24 hours, the reaction mixture was cooled, filtered through a celite pad and the resulting filtrate was then concentrated *in vacuo* to provide a viscous orange oil. Flash chromatography of this oil over silica gel with 60% ethyl acetate/hexane plus 0.5% ammonium hydroxide as eluent gave the protected analog of the title compound in 61% yield.

The above-mentioned protected analog (635 mg, 1.4 mmol) was dissolved in 10 ml of dry tetrahydrofuran and the resulting solution was chilled to -78°C. Once chilled, 1.5 ml (2.39 mmol) of a 1.7M solution of n-butyllithium in hexane was added dropwise via syringe. Once n-butyllithium addition was complete the reaction mixture was warmed to room temperature. The reaction mixture was quenched with a saturated NaHCO₃ solution and then partitioned between ethyl acetate and water. The aqueous layer was extracted with ethyl acetate, and the organic layers were combined, washed with a saturated brine solution, dried over sodium sulfate and then concentrated *in vacuo* to provide a viscous orange oil. This oil was chromatographed over silica gel (elution with 20% ethyl acetate/hexane plus 0.5% ammonium hydroxide) to provide 161 mg of title compound as a pale yellow oil.

MS m/e 324(FD)

[α]_D -45.63°(MeOH)

Analysis calculated for C ₂₁ H ₂₈ N ₂ O:			
Theory:	C, 77.74;	H, 8.70;	N, 8.63;
Found:	C, 78.74;	H, 8.82;	N, 8.27.

Example 14**Preparation of (+)(2aS,4R)-6-(2-furyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole**

The title compound was prepared substantially in accordance with the method set forth in Example 13, above, utilizing 1.5 g (3.07 mmol) of (-)(2aS,4R)-1-benzoyl-6-iodo-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole, 250 mg of bis(triphenylphosphine)palladium(II) chloride and 1.21 g (3.38 mmol) of 2-(tributylstannyl)furan to provide 592 mg of title compound as a viscous brown oil.

MS m/e 325.22(FD)

[α]_D +42.0°(MeOH)

Analysis calculated for C ₂₁ H ₂₈ N ₂ O:			
Theory:	C, 77.74;	H, 8.70;	N, 8.63;
Found:	C, 77.59;	H, 8.10;	N, 8.83.

Example 15**Preparation of (+)(2aS,4R)-6-(3-furyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole**

The title compound was prepared substantially in accordance with the method described in Example 13, above, utilizing 1.50 g (3.07 mmol) of (+)(2aS,4R)-1-benzoyl-6-iodo-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole, 1.21 g (3.38 mmol) of 3-(tributylstannyl)furan and 250 mg of bis(triphenylphosphine)palladium(II) chloride to provide 711 mg of title product as a pale yellow viscous oil.

MS m/e 324(FD)

Analysis calculated for C ₂₁ H ₂₈ N ₂ O:			
Theory:	C, 77.24;	H, 8.70;	N, 8.63;
Found:	C, 77.49;	H, 8.68;	N, 8.45.

Example 16**Preparation of (+)(2aS,4R)-6-(2-thienyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole**

The title compound was prepared substantially in accordance with the method set forth in Example 13, above, utilizing 1.5 g (3.1 mmol) of (-)(2aS,4R)-1-benzoyl-6-iodo-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole, 150 mg of bis(triphenylphosphine)palladium(II) chloride and 1.27 g (3.41 mmol) of 2-(tributylstannyl)thiophene to provide 719 mg of title compound as a light brown viscous oil.

MS m/e 341(FD)

Analysis calculated for C ₂₁ H ₂₈ N ₂ S:				
Theory:	C, 74.07;	H, 8.29;	N, 18.60;	S, 9.42;
Found:	C, 74.24;	H, 8.60;	N, 17.52;	S, 9.15.

Example 17**Preparation of (+)(2aS,4R)-6-(2-pyridyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole**

The title compound was prepared substantially in accordance with the method set forth in Example 13, above, utilizing 1.50 g (3.07 mmol) of (-)(2aS,4R)-1-benzoyl-6-iodo-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole, 250 mg of bis(triphenylphosphine)palladium(II) chloride and 1.24 g (3.38 mmol) of 2-(tributylstannyl)pyridine to produce 474 mg of title compound as a colorless foam. The hydrochloride salt of the title compound was prepared by dissolving the foam in diethyl ether and then treating the resulting solution with a saturated hydrochloric acid in methanol solution. A yellow foam comprised of such salt was afforded after concentration *in vacuo*.

MS m/e 336.24(FD)

Analysis calculated for $C_{22}H_{29}N_3 \cdot HCl$:

Theory:	C, 71.04;	H, 8.13;	N, 11.30;
Found:	C, 70.60;	H, 8.46;	N, 10.58.

Example 18

Preparation of (+)(2aS,4R)-6-(3-pyridyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

The title compound was prepared substantially in accordance with the procedure set forth in Example 13, above, utilizing 1.50 g (3.07 mmol) of (-)(2aS,4R)-1-benzoyl-6-iodo-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole, 250 mg of bis(triphenylphosphine)palladium(II) chloride and 1.24 g (3.38 mmol) of 3-(tributylstannyl)pyridine to produce 475 mg of title compound as a pale yellow oil. The dihydrochloride salt of the title compound was prepared by dissolving the oil in diethyl ether and then adding a saturated hydrochloric acid in methanol solution dropwise. Once an excess of hydrochloric acid had been added the mixture was concentrated *in vacuo* to provide a pale yellow foam.

MS m/e 336.24(FD)

Analysis calculated for $C_{22}H_{29}N_3 \cdot 2HCl$:

Theory:	C, 64.70;	H, 7.65;	N, 10.29;
Found:	C, 65.84;	H, 7.55;	N, 9.76.

Example 19

Preparation of (-)(2aR,4S)-6-(2-oxazolyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

A. 2-tributylstannyloxazole

A solution of 1.00 g (14.5 mmol) of oxazole in 25 ml of THF at -78°C was treated with 10.2 ml (14.6 mmol) of 1.43M butyllithium in hexane. After stirring for 30 minutes, an addition of 3.93 ml (14.5 mmol) of tributyltin chloride was made, and the solution was allowed to warm to room temperature. Stirring was continued for another hour after which most of the solvents were evaporated *in vacuo*. The resulting residue was taken up in 50 ml of hexane, and the resulting precipitate was separated by filtration through filtercel. Evaporation of the solvent from the filtrate provided 5.13 g of a colorless oil which was identified by NMR as the 2-stannyl derivative plus a small amount of tetrabutylstannane.

B. (-)(2aR,4S)-1-benzoyl-6-(2-oxazolyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

A solution of 5.0 g (13.8 mmol) of the crude 2-tributylstannyloxazole prepared above and 6.8 g (13.9 mmol) of (+)(2aR,4S)-1-benzoyl-6-iodo-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole in 100 ml of toluene was treated with 0.7 g (0.6 mmol) of tetrakis-(triphenylphosphine)palladium then refluxed under nitrogen for 20 hours. After cooling the reaction mixture was washed with a saturated brine solution and then dried over Na_2SO_4 . Concentration *in vacuo* provided a viscous oil which was chromatographed over a silica gel column using a solvent gradient progressing from toluene to 1:1 toluene/EtOAc. The product from the column was dissolved in 1M HCl. This solution was then washed with ether, made alkaline with 5M NaOH, and extracted with CH_2Cl_2 . Concentration of the extract *in vacuo* gave about 4 g of a brown oil. When this oil was dissolved in pentane a small amount of a red/brown resin separated leaving a clear, yellow solution. The resin was separated and the pentane was evaporated to provide a residue. This residue was crystallized by dissolving it in a small amount of CH_2Cl_2 and slowly adding isooctane. The crystalline (-)(2aR,4S)-1-benzoyl-6-(2-oxazolyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole.

obtained in four crops, weighed 2.63 g. m.p. 103-104°C.

C. (-)(2aR,4S)-6-(2-oxazoly)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

A solution of 1.00 g (2.33 mmol) of the above 1-benzoyl compound in 25 ml of THF was stirred at -78°C as 3.0 ml (4.29 mmol) of 1.43M butyllithium in hexane was added. The resulting solution was allowed to warm to 0°C, then poured into water and extracted with CH₂Cl₂. The CH₂Cl₂ extract was then, in turn, extracted with 1M HCl. The resulting aqueous extract was made alkaline with 1M NaOH, and, in turn, extracted with CH₂Cl₂. After drying over Na₂SO₄, the extract was concentrated in vacuo to provide title compound as a viscous oil. m.p. 103-104°C

MS m/e 326

[α]_D = -60° (C=1, MeOH).

Analysis calculated for C₂₀H₂₇N₃O:

Theory:	C, 73.81;	H, 8.36;	N, 12.91;
Found:	C, 73.37;	H, 8.26;	N, 12.09.

Example 20

Preparation of (-)(2aR,4S)-6-(5-isoxazoly)-4-[di(cyclopropylmethyl)amino]-1,2,2a,3,4,5-hexahydrobenz[cd]indole

To a solution of (-)(2aR,4S)-6-acetyl-4-[di(cyclopropylmethyl)amino]-1,2,2a,3,4,5-hexahydrobenz-[cd]indole (2.5 g, 7.7 mmol) and triethylamine (1.1 ml, 8 mmol) in 90 ml CH₂Cl₂ was added dropwise a solution of 2,2,2-trichloroethylchloroformate (1.7 g, 8 mmol) in 10 ml CH₂Cl₂. The reaction mixture was stirred at room temperature for one hour and then extracted with water and 1N HCl. The organic solution was washed with a saturated NaHCO₃ solution, and a saturated brine solution, dried over MgSO₄ and then concentrated to dryness in vacuo to give 3.1 g of the 1-carbamylindoline.

A solution of the 1-carbamylindoline (3.1 g, 6.2 mmol) and tris(dimethylamino)methane (5 ml) in 100 ml of toluene was stirred at reflux for 16 hours. After 16 hours the reaction mixture was concentrated to dryness in vacuo. The resulting residue was dissolved in 50 ml of acetic acid and 2.0 g (29 mmol) of hydroxylamine hydrochloride were added. The resulting reaction mixture was stirred at room temperature for 16 hours and then concentrated to dryness in vacuo. The resulting residue was suspended in water and an excess of a concentrated NH₄OH solution was added to basify the mixture. The basic mixture was then extracted with CH₂Cl₂ and the resulting organic extract was washed with a saturated brine solution, dried over MgSO₄ and then concentrated in vacuo to give 2.1 g of an oil. This oil was chromatographed (flash column, silica gel, EtOAc) to yield 1.7 g of the protected (+)(2aR,4S)-6-(5-isoxazoly)indoline.

The protected isoxazolyindoline (1.7 g, 3.2 mmol) was dissolved in 30 ml of acetic acid and 1.5 g of zinc dust were added all at once. The resulting reaction mixture was stirred at room temperature for four hours and then filtered through a celite pad. The filtrate thus obtained was then concentrated to dryness in vacuo. The resulting residue was suspended in a saturated NaHCO₃ solution then extracted with CH₂Cl₂. The organic extract was then washed with a saturated brine solution, dried over MgSO₄ and concentrated in vacuo to an oil. This oil was chromatographed (flash column, silica gel, EtOAc) to give 660 mg of title compound.

Example 21

Preparation of (-)(2aR,4S)-6-(5-oxazoly)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

A. (-)(2aR,4S)-6-bromo-1-trityl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

To a stirred solution of 12.8 g (29 mmol) of (+)(2aR,4S)-1-benzoyl-6-bromo-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole in 200 ml of tetrahydrofuran cooled to -78°C under a nitrogen atmosphere was added 20 ml (32 mmol) of a 1.6M solution of n-butyllithium in hexane. The reaction mixture was stirred at -78°C for 30 minutes and then allowed to warm to -20°C. To the reaction mixture was added 50 ml of a 1N hydrochloric acid solution. The mixture was extracted once with diethyl ether. The acidic solution was made alkaline with the addition of cold 5N sodium hydroxide solution. The basic mixture was extracted twice with methylene chloride. The combined organic solution was washed with a saturated sodium chloride solution. The methylene chloride solution was dried over magnesium sulfate and evaporated to give 9.8 g of (-)(2aR,4S)-6-bromo-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole.

To a solution of the above product (9.6 g, 0.028 mol) and triethylamine (3.03 g, 0.03 mol) in 100 ml of

methylene chloride was added a solution of trityl chloride (7.8 g, 0.028 mol) in 100 ml of methylene chloride dropwise at room temperature. The reaction mixture was stirred for 16 hours at room temperature. The reaction mixture was extracted with water and a cold 1N hydrochloric acid solution. The organic solution was washed with a saturated sodium bicarbonate solution and with a saturated brine solution. The organic solution was dried over magnesium sulfate and concentrated to dryness *in vacuo* to give a residue. The residue was slurried with warm hexane, cooled and filtered to remove insolubles. The filtrate was concentrated to an oil. The oil was chromatographed (silica gel, 20% ethyl acetate in hexane) to provide 12.7 g of the above-titled compound.

B. (-)(2aR,4S)-6-formyl-1-trityl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

To a solution of (-)(2aR,4S)-6-bromo-1-trityl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (6.8 g, 12 mmol) in 100 ml of tetrahydrofuran cooled to -78°C under a nitrogen atmosphere was added dropwise a 1.6M solution of n-butyllithium in hexane. The reaction mixture was stirred at -78°C for 1 hour. Dimethylformamide (3 ml) was added to the reaction mixture and the mixture was stirred at room temperature for 30 minutes. The reaction mixture was quenched with water and then extracted with ethyl acetate. The ethyl acetate solution was washed with a saturated brine solution, dried over magnesium sulfate and concentrated to dryness to provide 5.6 g of the above-titled compound as an oil.

C. (-)(2aR,4S)-6-(5-oxazolyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

A reaction mixture of (-)(2aR,4S)-6-formyl-1-trityl-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (1.06 g, 2 mmol), tosylmethyl isocyanide (390 mg, 2 mmol) and potassium carbonate (304 mg, 2.2 mmol) in 100 ml of methanol was stirred at reflux temperature under a nitrogen atmosphere for 16 hours. The reaction mixture was concentrated to dryness and water was added to the residue. The aqueous mixture was extracted with ethyl acetate. The ethyl acetate solution was washed with a saturated brine solution, dried over magnesium sulfate and concentrated to dryness to yield 1 g of an oil. The oil was dissolved in 20 ml of tetrahydrofuran and 50 ml of 5N hydrochloric acid solution. The reaction mixture was stirred at room temperature for 30 minutes. The reaction mixture was extracted twice with ethyl acetate and the acidic solution was then made alkaline by addition of excess concentrated ammonium hydroxide solution. The basic mixture was extracted twice with ethyl acetate. The ethyl acetate solution was washed with a saturated brine solution, dried over magnesium sulfate and concentrated to dryness to provide 0.5 g of an oil. The oil was purified by silica gel chromatography, with ethyl acetate as eluent, to give 0.3 g of title compound.

MS (FD) m/e/ 325

Example 22

Preparation of (2aR,4S)-6-(3-pyridyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

A. 3-pyridylboronic acid

A solution of 4.0 ml (6.56 g, 42 mmol) of 3-bromopyridine and 9 ml (8 mmol) of trimethylborate in 100 ml of diethyl ether was cooled to -70°C then treated slowly with 33 ml (83.8 mmol) of a 2.54M tert-butyllithium in pentane solution. After allowing the resulting slurry to warm to room temperature the solvents were evaporated under vacuum. The residual oil was treated carefully with 50 ml of a 1M hydrochloric acid solution. Several milliliters of methylene chloride were added and the mixture was stirred until the oil had dissolved. The aqueous layer was washed with fresh methylene chloride. The pH of the aqueous solution was raised to 12 with 5M sodium hydroxide solution and the washing was repeated. The pH of the aqueous solution was then lowered to 6.5 with concentrated hydrochloric acid solution. After chilling, this solution was filtered, saturated with sodium chloride, then extracted several times with a 2:1 mixture of diethyl ether and isopropanol. Evaporation of these extracts produced a colorless solid. This material was further purified by dissolving in methanol, evaporating to a thick paste, adding a few milliliters of water, concentrating further under vacuum, then chilling and collecting the crystalline product. Additional product in the aqueous mother liquor was isolated by repeating this process. Thorough drying of this hydrated product at 0.1 mm pressure afforded a fine powder weighing 2.2 g. Elemental analysis and a mass spectrum indicated the product so isolated was primarily the anhydride (tripyridylboroxane).

B. (2aR,4S)-1-benzoyl-6-(3-pyridyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

A solution of 4.00 g of (2aS,4R)-1-benzoyl-6-iodo-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole and 0.60 g (0.525 mmol) of tetrakis(triphenylphosphine)palladium in 50 ml of toluene was combined with 10 ml of a 2M sodium bicarbonate solution. This mixture was treated with 1.1 g (9.0 mmol) of the above 3-pyridylboronic anhydride, and it was stirred vigorously at 105°C under nitrogen for 24 hours. The half-complete reaction was charged with an additional 1.0 g of 3-pyridylboronic anhydride and heating was continued another 24 hours. The cooled mixture was filtered through filtercel. The organic layer was

washed with a saturated sodium chloride solution. The toluene was evaporated and the residue was partitioned between 1M hydrochloric acid solution and methylene chloride. The aqueous layer was basified with 5M sodium hydroxide solution and the product was extracted into methylene chloride. After washing with a saturated sodium chloride solution and drying over sodium sulfate the methylene chloride was evaporated leaving a viscous oil. This crude product was chromatographed over a silica gel column using 10% ethyl acetate in toluene, then 25% ethyl acetate in toluene, and finally 1:1 ethyl acetate/toluene as eluent. The purified (2aR,4S)-1-benzoyl-6-(3-pyridyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole was an oil weighing 2.99 g.

C. (2aR,4S)-6-(3-pyridyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

A solution of 2.75 g (6.26 mmol) of the above 1-benzoyl compound in 50 ml of tetrahydrofuran was stirred at -78°C as 7.7 ml (11.3 mmol) of a 1.47M butyllithium in hexane solution was added. This solution was allowed to warm to 0°C, then poured into water and extracted with methylene chloride. The methylene chloride was evaporated and the residue was chromatographed over 50 g of florisil using ethyl acetate as eluent. The product from the column was a pale yellow oil weighing 2.0 g which assayed as title product.

Example 23

Preparation of (-)(2aR,4S)-6-(3-isoxazoly)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

To a cool (-5°C) solution of 2.6 g (3.6 mmol) of (-)(2aR,4S)-1-triphenylmethyl-6-(1-oximidoethane)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (prepared substantially in accordance with the oxime substrate prepared in Example 8) in 100 ml of tetrahydrofuran was added 6.9 ml of a 1.6M solution of n-butyllithium in hexane. The resulting solution was stirred at -5°C for one hour and then 2 ml (26 mmol) of dimethylformamide was added all at once. The resulting solution was warmed to room temperature and then stirred for one more hour. After stirring at room temperature for one hour, the reaction solution was poured into 50 ml of a 1N sulfuric acid solution. The acidic solution was warmed on a steam bath for one hour, cooled to room temperature and then extracted with diethyl ether to remove impurities. The acidic solution was made alkaline with excess 5N sodium hydroxide solution and then extracted with ethyl acetate. The extract was washed with a saturated brine solution, dried over magnesium sulfate and then concentrated *in vacuo* to provide 1 g of an oil. This oil was purified by flash chromatography (using ethyl acetate as eluent) to provide 400 mg of the title compound as an oil.

Example 24

Preparation of (+)(4S)-6-(3-isoxazoly)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole.

A mixture of (+)(2aS,4R)-6-(3-isoxazoly)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (180 mg, 0.5 mmol) and 1 g of MnO₂ in 40 mL of CH₂Cl₂ was sonicated at 50-55 KHz for 2 hr. The reaction mixture warmed to reflux during the time period. The reaction mixture was filtered through a celite pad and the filtrate was concentrated to dryness *in vacuo*. The residue was chromatographed (flash column, silica gel, ethyl acetate) to provide 50 mg of the isoxazole indole product as an oil.

MS (FD) 323

Example 25

Preparation of (-)(4R)-6-(3-isoxazoly)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole

A mixture of (-)(2aR,4S)-6-(3-isoxazoly)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (400 mg, 1.2 mmol) and 1 g of MnO₂ in 100 ml of CH₂Cl₂ was sonicated at 50-55 KHz for four hours. The reaction mixture was warmed to reflux during the time period. After four hours, the reaction mixture was filtered through a celite pad and the filtrate was concentrated to dryness *in vacuo*. The resulting residue was chromatographed (flash column, silica gel, EtOAc) to provide 55 mg of title product as an oil. MS m/e 323 (FD)

NMR (300 MHz, CDCl₃) δ 0.90 (t, 6H); 1.2-1.6 (m, 5H); 2.2-3.6 (m, 8H); 6.6 (s, 1H); 6.9 (s, 1H); 7.2 (d, 1H); 7.4 (d, 1H); 8.0 (bs, 1H); 8.4 (s, 1H).

Example 26

Preparation of (-)(4R)-6-(5-isoxazoly)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole

A mixture of (-)(2aR,4S)-6-(5-isoxazoly)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (253 mg, 0.8 mmol) and 1 g of MnO₂ in 100 ml of CH₂Cl₂ was sonicated at 50-55 KHz for four hours. The reaction mixture was warmed to reflux during the time period. After four hours, the reaction mixture was filtered through

a celite pad and the filtrate was concentrated to dryness in vacuo. The resulting residue was chromatographed (flash column, silica gel, EtOAc) to provide 130 mg of title product. Recrystallization from hexane gave 60 mg of purified title product.

mp 126-127°C

5 MS m/e 323(FD)

Analysis calculated for $C_{20}H_{25}N_3O$:

Theory:	C, 74.27;	H, 7.79;	N, 12.99;
Found:	C, 74.44;	H, 7.71;	N, 12.78.

10

Example 27

15 Preparation of (-)-(4R)-6-(5-isoxazolyl)-4-[di(cyclopropylmethyl)amino]-1,3,4,5-tetrahydrobenz[cd]indole

A mixture of (-)-(2aR,4S)-6-(5-isoxazolyl)-4-[dicyclopropylmethyl]amino-1,2,2a,3,4,5-hexahydrobenz[cd]indole (660 mg, 1.9 mmol) and 3 g of MnO_2 in 100 ml of CH_2Cl_2 was sonicated at 50-55 KHz for four hours. The reaction mixture was warmed to reflux during the time period. After four hours, the reaction mixture was filtered through a celite pad and the filtrate was concentrated to dryness in vacuo. The resulting residue was chromatographed (flash column, silica gel, EtOAc) to provide 300 mg of title product. Recrystallization from hexane gave 125 mg of purified title product.

20

mp 146-147°C

MS m/e 347(FD)

25

$[\alpha]_D = -43.58^\circ$ (MeOH).

Analysis calculated for $C_{22}H_{25}N_3O$:

Theory:	C, 76.05;	H, 7.25;	N, 12.09;
Found:	C, 76.09;	H, 7.37;	N, 12.10.

30

Example 28

35 Preparation of (-)-(4R)-6-(3-pyrazolyl)-4-[di(cyclopropylmethyl)amino]-1,3,4,5-tetrahydrobenz[cd]indole

A solution of (-)-(4R)-6-acetyl-4-[di(cyclopropylmethyl)amino]-1,3,4,5-tetrahydrobenz[cd]indole (0.5 g, 1.6 mmol) and 2 ml of tris(dimethylamino)methane in 100 ml of toluene was refluxed for 16 hours. After 16 hours, the reaction solution was concentrated to dryness in vacuo and the resulting residue was dissolved in 100 ml of ethanol. Two milliliters of an 85% hydrazine solution were added to the ethanolic solution and the resulting reaction mixture was stirred at reflux temperature for 2 hours. After 2 hours, the reaction solution was, again, concentrated to dryness in vacuo. The resulting residue was then dissolved in 50 ml of 1N hydrochloric acid and the acidic solution was extracted with EtOAc. After extraction with EtOAc the acidic solution was then basified by adding an excess of a concentrated NH_4OH solution. The basic mixture was then extracted with EtOAc. The organic extract was washed with a saturated brine solution, dried over $MgSO_4$ and concentrated to dryness in vacuo to provide 500 mg of an oil. This oil was purified via chromatography (silica gel, flash column, EtOAc) to provide 400 mg of title compound as an oil.

45

MS m/e 346(FD)

NMR (300 MHz, $CDCl_3$) δ 0.1 (bs, 4H); 0.5 (m, 4H); 0.95 (m, 2H); 1.3 (dd, 1H); 2.6-3.8 (m, 9H); 6.4 (s, 1H); 6.9 (s, 1H); 7.2 (d, 1H); 7.3 (d, 1H); 7.6 (s, 1H); 8.2 (bs, 1H).

50

Example 29

Preparation of (-)-(4R)-6-(3-pyrazolyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-tetrahydrobenz[cd]indole

55 A solution of (-)-(4R)-6-acetyl-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole (0.4 g, 1.3 mmol) and 2 ml of tris(dimethylamino)methane in 100 ml of toluene was refluxed for 16 hours. After that time, the reaction solution was concentrated to dryness in vacuo and the resulting residue was dissolved in 100 ml of ethanol. Two milliliters of an 85% hydrazine solution were added to the ethanolic solution and the resulting reaction mixture was stirred at reflux for 2 hours. After 2 hours, the reaction solution was concentrated to dryness in

vacuo and the resulting residue was dissolved in 50 ml of 1N hydrochloric acid. The acidic solution was extracted with EtOAc and then basisified by adding an excess of a concentrated NH_4OH solution. The basic mixture was then extracted with EtOAc. The organic extract was washed with a saturated brine solution, dried over MgSO_4 and then concentrated *in vacuo* to provide an oil. This oil was purified by chromatography (silica gel, flash column, EtOAc) to provide 400 mg of title compound as an oil.

MS m/e 322(FD)

NMR (300 MHz CDCl_3) δ 0.9 (t, 6H); 1.5 (m, 4H); 2.4-2.6 (m, 4H); 2.8 (dd, 1H); 3.0 (m, 2H); 3.1-3.3 (m, 2H); 6.5 (s, 1H); 6.9 (s, 1H); 7.1 (d, 1H); 7.3 (bs, 1H); 7.7 (d, 1H); 8.1 (bs, 1H).

Example 30

Preparation of (-)(4R)-6-(3-phenyl-1,2,4-oxadiazol-5-yl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole

Magnesium dioxide (200 mg) was added to a solution of (-)(2aR,4S)-6-(3-phenyloxadiazol-5-yl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (40 mg, 0.10 mmol) in 15 ml of hexane. The resulting mixture was then sonicated at 50-55 KHz, while maintaining the reaction solution's temperature between 25-35°C, for 135 minutes. After that time the reaction mixture was filtered through celite and the recovered solid was washed sequentially with hexane and CH_2Cl_2 . The filtrate was combined with the organic washes and the resulting solution was washed sequentially with water and a saturated brine solution, dried over Na_2SO_4 and then concentrated *in vacuo* to give 30 mg of an orange film. This film was purified by flash chromatography [2:3 ether:hexane (NH_4OH)] to give 70 mg of title product. This product was purified by recrystallization from a CH_2Cl_2 /hexane solvent system to give 50 mg of title compound as a light green solid.

mp 154-155°C

Analysis calculated for $\text{C}_{25}\text{H}_{28}\text{N}_4\text{O} \cdot 0.25 \text{H}_2\text{O}$:

Theory:	C, 74.14;	H, 7.09;	N, 13.83;
Found:	C, 74.21;	H, 7.01;	N, 13.58.

Example 31

Preparation of (-)(4R)-6-(2-oxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole

A solution of 0.65 g (2.0 mmol) of (-)(2aR,4S)-6-(2-oxazolyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole in 30 ml of CH_2Cl_2 was sonicated (50-55 KHz) in the presence of 2.5 g of MnO_2 for 5 hours. The oxidant was removed by filtration through celite. The crude product obtained after filtering and removing the solvent was chromatographed over 15 g of silica gel using 1:9 EtOAc/toluene, then crystallized from isooctane to provide 0.24 g of title compound.

mp 73-74°C

Analysis calculated for $\text{C}_{20}\text{H}_{25}\text{N}_3\text{O}$:

Theory:	C, 74.27;	H, 7.79;	N, 12.99;
Found:	C, 73.97;	H, 7.84;	N, 12.90.

NMR (300 MHz, CDCl_3) δ 0.90 (t, 6H); 1.49 (sextet, 4H); 2.60 (t, 4H); 2.82 (t, 1H); 3.00 (m, 1H); 3.00 (dd, 1H); 3.26 (m, 1H); 3.78 (m, 1H); 6.90 (s, 1H); 7.28 (s, 1H); 7.73 (s, 1H); 7.22 (d, 1H); 7.86 (d, 1H); 7.97 (s, 1H).

MS m/e 324(FD)

$[\alpha]_D = -60^\circ$ (c=1, MeOH)

Example 32

Preparation of (-)(4R)-6-(5-oxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole

A mixture of (-)(2aR,4S)-6-(5-oxazolyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole (300 mg, 0.9 mmol) and 0.5 g of MnO_2 in 50 ml of methylene chloride was stirred at room temperature for 16 hours. The oxidant was removed by filtration through celite. The crude product obtained after filtering and removing the solvent was purified by flash chromatography, using ethyl acetate as eluent, to provide 100 mg of title com-

pound. Recrystallization from hexane gave 60 mg of pure title compound.

m.p. 161-162°C

MS (FD) 323

$[\alpha]_D -64.0^\circ$ (MeOH)

Analysis calculated for $C_{20}H_{25}N_3O$:

Theory:	C, 74.27;	H, 7.79;	N, 12.99;
Found:	C, 73.99;	H, 7.82;	N, 12.74.

Example 33

Preparation of (-)(4R)-6-(3-pyridyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole

A solution of 1.0 g (2.99 mmol) of (2aR,4S)-6-(3-pyridyl)-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole and 1.0 g (9 mmol) of indole in 50 ml of THF was treated at 0°C with 0.61 g (1.70 mmol) of benzeneseleninic anhydride. The resulting solution was allowed to warm to room temperature. After 2 hours the reaction solution was diluted with water and the product was extracted into methylene chloride. The methylene chloride was evaporated and the resulting residue was taken up in aqueous tartaric acid. This acidic solution was washed with methylene chloride, then basified with 5M sodium hydroxide solution. The product was extracted into methylene chloride. After evaporating the methylene chloride the product was chromatographed over 25 g of silica gel using 10% ethyl acetate in toluene followed by 1:1 ethyl acetate/toluene as eluent. The material from the column was crystallized from toluene/hexane to afford 0.53 g of title compound.

m.p. 185-186°C

MS (FD) 334

$[\alpha]_D -52.0^\circ$ (MeOH)

NMR (300 MHz, $CDCl_3$) δ 0.88 (t,6H), 1.45 (sextet,4H), 2.52 (t,4H), 2.45 (dd,1H), 3.03 (m,3H), 3.17 (m,1H), 6.94 (s,1H), 7.17 (d,1H), 7.27 (d,1H), 7.38 (dd,1H), 7.77 (d,1H), 8.01 (s,1H), 8.57 (d,1H), 8.74 (s,1H).

Analysis calculated for $C_{22}H_{27}N_3$:

Theory:	C, 79.24;	H, 8.16;	N, 12.60;
Found:	C, 79.07;	H, 8.15;	N, 12.78.

Example 34

Preparation of (4R)-6-[2-(1,3,4-oxadiazolyl)]-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole

To a solution of 5.0 g (13.1 mmol) of (4S)-6-iodo-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole in 250 ml of acetonitrile was added 3.5 g (37 mmol) of 2-hydroxypyridine, 16.9 g (122 mmol) of powdered potassium carbonate, and 0.30 g (0.26 mmol) of tetrakis(triphenylphosphine)palladium. The resultant mixture was heated at 65°C under a carbon monoxide atmosphere for 14 hours. The cooled mixture was then treated with 2 ml of hydrazine hydrate and stirred for another 15 hours. After filtering, the mixture was concentrated *in vacuo* to provide a residue. This residue was taken up in methylene chloride and extracted with aqueous tartaric acid solution. The aqueous layer was basified with 1M sodium hydroxide solution and the product was extracted into methylene chloride. Evaporation of solvent *in vacuo* left an oily solid which, after thorough washing with a 1:2 mixture of toluene and hexane, afforded 0.60 g of (4R)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole-6-carboxylic hydrazide. m.p. 199-201°C

A solution of 0.20 g (0.64 mmol) of the above hydrazide in 10 ml of triethylorthoformate was heated at 150°C under nitrogen for 15 hours. Most of the excess triethylorthoformate was evaporated *in vacuo*. The resulting residue was dissolved in aqueous tartaric acid and washed with methylene chloride. After basifying the aqueous solution with sodium carbonate the product was extracted into methylene chloride. Evaporation of solvent *in vacuo* left crystalline product which was then chromatographed over 3 g of silica gel using ethyl acetate as eluent. The product from the column was recrystallized from toluene/hexane giving 0.16 g of title compound.

m.p. 137-138.5°C

MS (FD) 324

$[\alpha]_D -93.0^\circ$ (MeOH)

NMR (300 MHz, DMSO- d_6) δ 0.86 (t,6H), 1.40 (sextet,4H), 2.52 (qt,4H), 2.74 (dd,1H), 2.88 (dd,1H), 3.00 (dd,1H), 3.11 (m,1H), 3.63 (d,1H), 7.07 (s,1H), 7.26 (d,1H), 7.60 (d,1H), 9.23 (s,1H).

Analysis calculated for $C_{19}H_{24}N_4O$:

Theory:	C, 70.34;	H, 7.46;	N, 17.27;
Found:	C, 70.13;	H, 7.33;	N, 16.98.

Example 35

Preparation of (-)(4R)-6-(5-phenyl-1,3,4-oxadiazol-2-yl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole

A solution of 0.20 g (0.64 mmol) of (4R)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole-6-carboxylic hydrazide in 2.0 ml of trimethylorthobenzoate was heated at 135°C under nitrogen for 5 hours. After cooling, the semisolid mixture was dissolved in methylene chloride and the product was extracted into aqueous tartaric acid. The aqueous solution was basified with sodium carbonate and the product was extracted into methylene chloride. Evaporation of the methylene chloride left crystalline product which was then chromatographed over 3 g of silica gel using ethyl acetate as eluent. The product from the column was recrystallized from toluene to give 0.21 g of title compound.

m.p. 215-216°C

MS (FD) 400

$[\alpha]_D -23.0^\circ$ (MeOH)

NMR (300 MHz, DMSO- d_6) δ 0.88 (t,6H), 1.41 (sextet,4H), 2.55 (t,4H), 2.74 (dd,1H), 2.82 (d,1H), 3.11 (m,2H), 3.70 (d,1H), 7.09 (s,1H), 7.30 (d,1H), 7.62 (m,3H), 7.77 (d,1H), 8.08 (m,2H).

Analysis calculated for $C_{25}H_{28}N_4O$:

Theory:	C, 74.97;	H, 7.05;	N, 13.99;
Found:	C, 74.70;	H, 7.15;	N, 13.89.

Example 36

Preparation of (4R)-6-(2-oxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole

A. (2aS,4R)-6-iodo-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole

A mixture of 100 g (0.21 mol) of (2aS,4R)-1-benzoyl-6-iodo-4-(di-n-propylamino)-1,2,2a,3,4,5-hexahydrobenz[cd]indole and 1 liter of 1M ethanolic potassium hydroxide was refluxed under nitrogen for 5 hours. The cooled mixture was diluted with two volumes of water, and the product was extracted into methylene chloride. After drying the extract over sodium sulfate the solvent was evaporated leaving 78 g of the debenzoylated product as a light brown oil.

B. (4S)-6-iodo-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole

A solution of 78 g (0.20 mol) of the above compound in 1 liter of methylene chloride was sonicated (50-55 KHz) in the presence of 100 g of MnO_2 for 24 hours. The oxidant was removed by filtration through filterrel. The crude product obtained after filtering and removing the solvent was chromatographed over silica gel using 3% ethyl acetate in toluene as eluent. The purified indole was a viscous, amber oil weighing 72 g.

C. (4R)-6-(2-oxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole

A suspension of 11.0 g (55 mmol) of potassium hydride (20% dispersion in oil) in 500 ml of tetrahydrofuran was maintained at 0°C as 20.0 g (52.4 mmol) of the above compound in a few milliliters of tetrahydrofuran was added. After stirring for 40 minutes, the cold solution was treated with 16.0 ml (60.4 mmol) of triisopropylsilyl triflate. Stirring was continued for another 2 hours. The mixture was then poured into cold sodium bicarbonate solution and the product was extracted into methylene chloride. The extract was washed with a saturated sodium chloride solution and then dried over sodium sulfate. Evaporation of solvent *in vacuo* left a brown oil which was chromatographed over silica gel using hexane followed by toluene as eluent. The silylated product from the column was isolated as a light brown oil weighing 26.4 g.

This oil, along with 25 g (69 mmol) of 2-tributylstannyloxazole, was dissolved in 500 ml of acetonitrile. The resulting solution was treated with 1.0 g of tetrakis(triphenylphosphine)palladium and then refluxed under nitrogen for 24 hours. Most of the acetonitrile was evaporated *in vacuo* to provide an oil. This oil was taken up in 200 ml of diethyl ether and washed with a 1% hydrogen peroxide solution. The organic layer was filtered, washed with a saturated sodium chloride solution and then dried over sodium sulfate. The solvent was evaporated and the residual oil was chromatographed over silica gel using toluene followed by ethyl acetate as eluent. The resulting coupled product (24 g) was dissolved in 250 ml of tetrahydrofuran. To this solution was added 5 g of boric acid. The solution was then cooled to 0°C and treated with 75 ml of a 1M tetrabutylammonium fluoride in tetrahydrofuran solution. After one hour the solution was diluted with water and the product was extracted into methylene chloride. The methylene chloride was then evaporated, diethyl ether was added and the product was extracted into aqueous tartaric acid (10 g in 500 ml). The aqueous solution was basified with 1M sodium hydroxide solution and the product was extracted into methylene chloride. After washing with a saturated sodium chloride solution and drying over sodium sulfate the methylene chloride was evaporated and the product was chromatographed over silica gel using toluene followed by 5% ethylacetate in toluene as eluent. Trituration of the chromatographed product with isooctane afforded 6.30 g of crystalline title product. m.p. 77-79°C

Example 37

20 Preparation of (-)-(4R)-6-(5-oxazolyl)-4-[di(cyclopropylmethyl)amino]-1,3,4,5-tetrahydrobenz[cd]indole

A. (2aR,4S)-6-(5-oxazolyl)-4-[di-(cyclopropylmethyl)amino]-1,2,2a,3,4,5-hexahydrobenz[cd]indole
A reaction solution of 4.4 g (8.0 mmol) of (2aR, 4S)-6-formyl-1-trityl-4-[di-(cyclopropylmethyl)amino]-1,2,2a,3,4,5-hexahydrobenz[cd]indole, 1.56 g (8.0 mmol) of tosylmethyl isocyanide and 1.1 g (8.0 mmol) of potassium carbonate in 200 ml of methanol was heated to reflux and stirred at that temperature for 16 hours. After 16 hours, the reaction solution was concentrated to dryness *in vacuo*. The resulting residue was dissolved in water and the resulting aqueous solution was extracted with ethyl acetate. The ethyl acetate extract was washed with a saturated brine solution, dried over magnesium sulfate and then concentrated to dryness *in vacuo* to provide an oil. This oil was dissolved in 150 ml of a 2/1 tetrahydrofuran/5N hydrochloric acid solution. Once all of the oil had dissolved, the resulting solution was stirred at room temperature for 30 minutes and then extracted twice with ethyl acetate. After extraction, the acidic solution was made alkaline by adding an excess of ammonium hydroxide. This basic solution was then extracted twice with ethyl acetate. These ethyl acetate extracts were then combined, washed with a saturated brine solution, dried over magnesium sulfate and then concentrated to dryness to provide 2.0 g of crude title compound as an oil. This oil was purified by flash chromatography (ethyl acetate as eluent) to provide 1.1 g of purified title compound as an oil.

MS(FD) - 349

30 B. (-)-(4R)-6-(5-oxazolyl)-4-[di-(cyclopropylmethyl)amino]-1,3,4,5-tetrahydrobenz[cd]indole

To a cold (0°C) solution of 1.1 g (3.0 mmol) of the compound prepared in (A) above and 1.05 g of indole in 50 ml of tetrahydrofuran were added 720 mg (2.0 mmol) of benzeneseleninic anhydride. The resulting solution was stirred at room temperature for 3 hours and then poured into a saturated sodium hydrogen carbonate solution. The resulting solution was extracted with ethyl acetate. The ethyl acetate extract was, in turn, extracted with 2M tartaric acid. The tartaric acid extract was made alkaline by adding an excess of concentrated ammonium hydroxide and the resulting basic solution was then extracted with ethyl acetate. This extract was washed with a saturated brine solution, dried over magnesium sulfate and then concentrated to dryness *in vacuo* to provide 1.0 g of crude title compound as a residue. This residue was purified by flash chromatography (ethyl acetate as eluent) to provide 800 mg of title compound. Recrystallization of such material from ethyl acetate/hexane gave 500 mg of purified title compound.

m.p. 135-136°C

MS(FD) 347

Analysis calculated for $C_{22}H_{26}N_3O$:

Theory:	C, 76.05;	H, 7.25;	N, 12.10
Found:	C, 75.91;	H, 7.41;	N, 11.98

Optical Rotation:

$[\alpha]_D$ (MeOH) = -64.36

The compounds of Formula 1 have been found to have selective affinity for the 5HT receptors in the brain with much less affinity for other receptors. Because of their ability to selectively bind to 5-HT receptors, the compounds of Formula 1 are useful in treating disease states which require alteration of 5-HT receptor function, particularly 5-HT_{1A}, and/or 5HT_{1D} but without the side effects which may be associated with less selective compounds. This alteration may involve reproducing (an agonist) or inhibiting (an antagonist) the function of serotonin. These disease states include anxiety, depression, excessive gastric acid secretion, motion sickness, hypertension, emesis, nausea, sexual dysfunction, cognition, senile dementia, migraine, consumptive disorders such as appetite disorders, alcoholism and smoking. The foregoing conditions are treated with a pharmaceutically effective amount of a compound of Formula 1 or a pharmaceutically acceptable salt thereof.

The term "pharmaceutically effective amount", as used herein, represents an amount of a compound of the invention which is capable of diminishing the adverse symptoms of the particular disease (for example motion sickness or emesis). The particular dose of compound administered according to this invention shall, of course, be determined by the particular circumstances surrounding the case, including the compound administered, the route of administration, the particular condition being treated, and similar considerations. The compounds can be administered by a variety of routes including the oral, rectal, transdermal, subcutaneous, intravenous, intramuscular or intranasal routes. A typical single dose for prophylactic treatment, however, will contain from about 0.01 mg/kg to about 50 mg/kg of the active compound of this invention when administered orally. Preferred oral doses will be about 0.01 to about 3.0 mg/kg, ideally about 0.01 to about 0.1 mg/kg. When a present compound is given orally it may be necessary to administer the compound more than once each day, for example about every eight hours. For IV administration by bolus, the dose will be from about 10 µg/kg to about 300 µg/kg, preferably about 20 µg/kg to about 50 µg/kg.

The following experiments were conducted to demonstrate the ability of the compounds of the Formula 1 to bind to 5-HT receptors. Such experiments demonstrate the utility of the compounds of Formula 1 in treating disease states (such as emesis and motion sickness) which require alteration of 5-HT receptor function.

The affinities of certain of the compounds of Formula 1 at the central 5-HT_{1A} receptors were determined using a modification of the binding assay described by Taylor, et al., *J. Pharmacol. Exp. Ther.*, 236, 118-125 (1988). Membranes for the binding assay were prepared from male Sprague-Dawley rats (150-250 g). The animals were killed by decapitation, and the brains were rapidly chilled and dissected to obtain the hippocampi. Membranes from the hippocampi were either prepared that day, or the hippocampi were stored frozen (-70°C) until the day of preparation. The membranes were prepared by homogenizing the tissue in 40 volumes of ice-cold Tris-HCl buffer (50 mM, pH 7.4 at 22°C) using a Techmar Tissumizer (setting 65 for 15 seconds), and the homogenate was centrifuged at 39800xg for 10 minutes. The resulting pellet was then resuspended in the same buffer, and the centrifugation and resuspension process was repeated three additional times to wash the membranes. Between the second and third washes the resuspended membranes were incubated for 10 minutes at 37°C to facilitate the removal of endogenous ligands. The final pellet was resuspended in 67 mM Tris-HCl, pH 7.4, to a concentration of 2 mg of tissue original wet weight/200 µl. This homogenate was stored frozen (-70°C) until the day of the binding assay. Each tube for the binding assay had a final volume of 800 µl and contained the following: Tris-HCl (50 mM), pargyline (10 µM), CaCl₂ (3 mM), [³H]8-OH-DPAT (1.0 nM), appropriate dilutions of the drugs of interest, and membrane resuspension equivalent to 2 mg of original tissue wet weight, for a final pH of 7.4. The assay tubes were incubated for 10 minutes at 37°C, and the contents were then rapidly filtered through GF/B filters (pretreated with 0.5% polyethylenimine), followed by four 1 ml washes with ice-cold buffer. The radioactivity trapped by the filters was quantitated by liquid scintillation spectrometry, and specific [³H]8-OH-DPAT binding to the 5-HT_{1A} sites was defined as the difference between [³H]8-OH-DPAT bound in the presence and absence of 10 µM 5-HT.

The results of the evaluation of various compounds of Formula 1 in the test system described above are set forth in Table 1, below. In Table 1, the first column provides the example number of the compound evaluated while the second column provides the amount of test compound (expressed in nanomolar concentration) required to inhibit the binding of [³H]8-OH-DPAT by 50% (indicated as IC₅₀).

TABLE 1

IN VITRO BINDING AT THE 5-HT _{1A} RECEPTOR	
EXAMPLE No.	5-HT _{1A} <i>in vitro</i> binding (IC ₅₀ , nM)
24	0.61
26	0.10
30	7.1
31	2.77
32	0.53
33	0.87
34	0.75
35	19.04

The affinities of certain of the compounds of Formula 1 at the central 5-HT_{1D} binding sites were determined using a modification of the binding assay described by Heuring and Peroutka, *J. Neurosci.*, 7, 894 (1987). Bovine brains were obtained and the caudate nuclei were dissected out and frozen at -70°C until the time that the membranes were prepared for the binding assays. At that time the tissues were homogenized in 40 volumes of ice-cold Tris-HCl buffer (50 mM, pH 7.4 at 22°C) with a Techmar Tissumizer (setting 65 for 15 seconds), and the homogenate was centrifuged at 39,800xg for 10 minutes. The resulting pellet was then resuspended in the same buffer, and the centrifugation and resuspension process was repeated three additional times to wash the membranes. Between the second and third washes the resuspended membranes were incubated for 10 minutes at 37°C to facilitate the removal of endogenous 5-HT. The final pellet was resuspended in Tris buffer to a concentration of 25 mg of original tissue wet weight/ml for use in the binding assay. Each tube for the binding assay had a final volume of 800 µl and contained the following: Tris-HCl (50 mM), pargyline (10 µM), ascorbate (5.7 mM), CaCl₂ (3 mM), 8-OH-DPAT (100 nM to mask 5-HT_{1A} receptors), mesulergine (100 nM to mask 5-HT_{1C} receptors), [³H]5-HT (1.7-1.9 nM), appropriate dilutions of the drugs of interest, and membrane resuspension equivalent to 5 mg of original tissue wet weight, for a final pH of 7.4. The assay tubes were incubated for 10 minutes at 37°C, and the contents were then rapidly filtered through GF/B filters (pretreated with 0.5% polyethylenimine), followed by four 1 ml washes with ice-cold buffer. The radioactivity trapped by the filters was quantitated by liquid scintillation spectrometry, and specific [³H]5-HT binding to the 5-HT_{1D} sites was defined as the difference between [³H]5-HT bound in the presence and absence of 10 µM 5-HT.

The results of the evaluation of various compounds of Formula 1 in the test system described above are set forth in Table 2 below. In Table 2, the first column provides the example number of the compound evaluated while the second column provides the amount of test compound (expressed in nanomolar concentration) required to inhibit the binding of [³H]5-HT by 50% (indicated as IC₅₀).

TABLE 2

In Vitro Binding Activity At The 5-HT _{1D} Receptor	
EXAMPLE No.	5-HT _{1D} <i>in vitro</i> binding (IC ₅₀ , nM)
24	18.60
26	46.36
31	280
32	74.45
33	29.4
34	577

In another experiment certain compounds of Formula 1 were evaluated to determine ability to affect serotonin turnover *in vivo*. As a measure of such affect on serotonin turnover, the decrease of the serotonin metabolite 5-hydroxyindole acetic acid (5HIAA) was measured. The following protocol was employed.

Male albino rats were injected either subcutaneously or orally with an aqueous solution of the compound tested. The pH of the solution was adjusted as necessary to solubilize the compound. A control of the solution without the test compound was similarly administered to a control animal. The rats were decapitated one hour later. The whole brain was then removed and frozen on dry ice for storage prior to analysis. 5-Hydroxyindole acetic acid (5HIAA) concentration in whole brain was measured by liquid chromatography with electrochemical detection as reported by Fuller and Perry, "Effects of Buspirone and its Metabolite, 1-(2-pyrimidinyl)piperazine, on Brain Monoamines and Their Metabolites in Rats", *J. Pharmacol. Exp. Ther.*, 248, p. 50-56 (1989), incorporated herein by reference. The results of this procedure are provided in Table 3.

In Table 3, Column 1 provides the Example No. of the compound tested while Column 2 provides the minimum effective dose (in mg/kg) required to produce a statistically significant decrease in 5HIAA concentration relative to control (generally a 15-20% decrease) for both oral and subcutaneous methods of administration.

TABLE 3

Effect On Serotonin Turnover In Vivo		
EXAMPLE No.	Minimum Effective Dose (mg/kg)	
	s.c.	p.o.
31	1.0	1.0
33	0.3	>10
34	0.1	10
35	>1.0	

The activity of certain of the compounds of Formula 1 on emesis and motion sickness was also evaluated directly in the following test systems.

Cat Assay

Adult female cats of mixed strains were obtained and housed such that they had free access to food and water except during the time of testing. Cats were selected for these studies based upon a minimum of 2 emetic episodes in 5 tests in response to a 30 minute rotation (0.28 Hz, 17 rpm) on a motion device such as that described in Crampton, et al., *Aviat. Space Environ. Med.*, 56, pp. 462-465 (1985). Single emetic response tests were conducted at intervals of at least two weeks to prevent habituation to the motion stimulus. Baseline responses (occurrence of vomiting) to motion following saline pretreatment were determined before and after the evaluation of a test compound. Subjects received subcutaneous injections of a test compound in sterile saline to an injection volume of 0.1 mg/kg or vehicle 30 minutes before motion testing. The results of such testing are reported in Table 4 below.

Table 4

Suppression of Motion Sickness in Cats			
Treatment	Dose of Test Compound ($\mu\text{g/kg}$)	Percent of Cats Vomiting	Change in Test Subject's Behavior Observed
Saline	-	70	No
Example No. 32	10	70	No
Example No. 32	20	50	No
Example No. 32	40	0	No
Saline	-	60	No
Example No. 37	40	50	No
Example No. 37	160	30	No
Example No. 37	320	10	No

Pigeon Assays

A group of male White Carneaux pigeons, weighing approximately 460 to 650 g (85-90% of their free-feeding body weights) were also tested. Animals were given free access to water and oyster shell grit except during the test sessions and were fed approximately 20 g of grain-based feed (Purina Pigeon Checkers) once per day. The colony room was illuminated from 6 a.m. to 6 p.m. daily. All testing was conducted during the illuminated phase.

The pigeons were first fed 20 grams of Purina Pigeon Checkers in their home cages. Five minutes later, the birds were weighed and injected intravenously with 10 mg/kg of cisplatin (cis-platinum II diammine dichloride; Sigma Chemical Co., ST. Louis, MO) and then placed in a plexiglas observation cage. After 45 minutes, either vehicle or the desired amount of test compound were administered by intramuscular (pectoral muscle) injection. The test compound was dissolved in distilled water with the aid of a few drops of lactic acid. The animals were then observed for the next 4.5 hours in order to determine the number of pigeons vomiting. The results of such testing are shown in the Table 5 below.

Table 5

Suppression of Cisplatin Induced Emesis in Pigeons			
Treatment	Dose of Test Compound (mg/kg)	# of Pigeons	Percent of Pigeons Vomiting
Vehicle	-	4	100
Example No. 32	0.64	4	50
Example No. 32	1.28	4	25

Another group of male White Carneaux pigeons, weighing approximately 460 to 650 g (85-90% of their free-feeding body weights) were also tested using ditolylguanidine (DTG) in place of cisplatin to induce emesis. Animals were given free access to water and oyster shell grit except during the test sessions and were fed approximately 20 g of grain-based feed (Purina Pigeon Checkers) once per day. The colony room was illuminated from 6 a.m. to 6 p.m. daily. All testing was conducted during the illuminated phase.

The pigeons were first fed 20 grams of Purina Pigeon Checkers in their home cages. Five minutes later, the birds were weighed, injected with various doses of test compound and returned to their home cages. The test compound was dissolved in distilled water with the aid of a few drops of lactic acid. All injections were given

into the pectoral muscle in a volume of 1 ml/kg. After 15 minutes, a 5.6 mg/kg dose of DTG was administered and the pigeons were placed into Plexiglas observation chambers. One hour later, the birds were removed from the observation chambers and returned to their home cages and the floor of the observation cage was examined for the presence of expelled food. The dependent variable in the instant study was the percent of birds at each dose that exhibited evidence of expelled food. The results of such study are presented in Table 6 below.

Table 6

Suppression of Ditolylguanidine Induced Emesis in Pigeons			
Treatment	Dose of Test Compound (mg/kg)	No. of Pigeons	Percent of Pigeons Vomiting
Vehicle	-	4	100
Example No. 32	0.04	4	100
Example No. 32	0.08	4	50
Example No. 32	0.16	4	50
Example No. 32	0.32	4	0
Example No. 31	0.08	4	100
Example No. 31	0.16	4	75
Example No. 31	0.32	4	25
Example No. 31	0.64	4	25
Example No. 31	1.28	4	0
Example No. 37	0.16	4	100
Example No. 37	0.32	4	0
Example No. 37	0.64	4	0

The data in Tables 1 through 6 establish that the compounds of Formula 1 can be used to treat emesis and motion sickness. The term "emesis", as used for purposes of the present invention, means vomiting (the actual expulsion of stomach contents) and retching (vomiting movements without expulsion of matter). Accordingly, the compounds of Formula 1 can be used to suppress emetic responses to provocative motion (motion sickness) or various chemical stimuli such as oncolytic agents (e.g., cisplatin) or other psychoactive agents (e.g., xylazine, analgesics, anesthetics and dopaminergic agents) and the like.

The method of the present invention encompasses treatment of emesis or motion sickness in a prophylactic manner (i.e., using the compounds of Formula 1 to prevent emesis or motion sickness in a mammal susceptible to such conditions before the conditions actually occur or re-occur). Such prophylactic method of administration may be especially appropriate in cases where the patient is susceptible to motion sickness and is about to go on a boat, car or plane trip which, normally, would result in the patient's suffering a motion sickness attack or the patient is about to undergo treatment with various chemical stimuli (cancer chemo and radiation therapy, analgesic and anesthetic agents, etc.) known to cause emesis.

The compounds of the present invention, and the compounds used in the method of the present invention, are preferably formulated prior to administration. Therefore, yet another embodiment of the present invention is a pharmaceutical formulation comprising a compound of Formula 1, or a pharmaceutically acceptable salt thereof, admixed with one or more pharmaceutically acceptable carriers, diluents or excipients therefor.

The present pharmaceutical formulations are prepared by known procedures using well known and readily available ingredients. In making the compositions of the present invention, the active ingredient will usually be mixed with an excipient, diluted by an excipient or enclosed within an excipient serving as a carrier which can be in the form of a capsule, sachet, paper or other container. When the excipient serves as a diluent, it can be a solid, semi-solid or liquid material which acts as a vehicle, carrier or medium for the active ingredient. Thus, the compositions can be in the form of tablets, pills, powders, lozenges, sachets, cachets, elixirs, sus-

pensions, emulsions, solutions, syrups, aerosols (as a solid or in a liquid medium), ointments containing for example up to 10% by weight of the active compound, soft and hard gelatin capsules, suppositories, sterile injectable solutions and sterile packaged powders.

Some examples of suitable excipients include lactose, dextrose, sucrose, sorbitol, mannitol, starches, gum acacia, calcium phosphate, alginates, tragacanth, gelatin, calcium silicate, microcrystalline cellulose, polyvinylpyrrolidone, cellulose, water, syrup, and methyl cellulose. The formulations can additionally include lubricating agents such as talc, magnesium stearate and mineral oil, wetting agents, emulsifying and suspending agents, preserving agents such as methyl and propylhydroxybenzoates, sweetening agents or flavoring agents. The compositions of the invention may be formulated so as to provide quick, sustained or delayed release of the active ingredient after administration to the patient by employing procedures well known in the art.

The compositions are preferably formulated in a unit dosage form, each dosage containing from about 0.5 to about 50 mg, more usually about 1 to about 10 mg of the active ingredient. The term "unit dosage form" refers to physically discrete units suitable as unitary dosages for human subjects and other mammals, each unit containing a predetermined quantity of active material calculated to produce the desired therapeutic effect, in association with a suitable pharmaceutical excipient.

The following formulation examples are illustrative only and are not intended to limit the scope of the invention in any way.

20 Formulation 1

Hard gelatin capsules suitable for use in treating motion sickness are prepared using the following ingredients:

	Quantity (mg/capsule)
(-)-6-(2-oxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole	25
Starch, dried	425
Magnesium stearate	10
Total	460 mg

The above ingredients are mixed and filled into hard gelatin capsules in 460 mg quantities.

35 Formulation 2

A tablet formula is prepared using the ingredients below:

	Quantity (mg/tablet)
(±)-6-[5-(2-methyltetrazolyl)]-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole	25
Cellulose, microcrystalline	625
Colloidal silicon dioxide	10
Stearic acid	5

The components are blended and compressed to form tablets each weighing 665 mg.

50 Formulation 3

A dry powder inhaler formulation suitable for treating emesis is prepared containing the following components:

	Weight %
(±)-6-(5-isoxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole	5
Lactose	95

The active compound is mixed with the lactose and the mixture added to a dry powder inhaling appliance.

Formulation 4

Tablets suitable for treating emesis each containing 60 mg of active ingredient are made up as follows:

(+)-6-(2-pyrazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole	60 mg
Starch	45 mg
Microcrystalline cellulose	35 mg
Polyvinylpyrrolidone (as 10% solution in water)	4 mg
Sodium carboxymethyl starch	4.5 mg
Magnesium stearate	0.5 mg
Talc	1 mg
Total	150 mg

The active ingredient, starch and cellulose are passed through a No. 20 mesh U.S. sieve and mixed thoroughly. The solution of polyvinylpyrrolidone is mixed with the resultant powders which are then passed through a No. 4 mesh U.S. sieve. The granules so produced are dried at 50-60°C and passed through a No. 16 mesh U.S. sieve. The sodium carboxymethyl starch, magnesium stearate and talc, previously passed through a No. 30 mesh U.S. sieve, are then added to the granules which, after mixing, are compressed on a tablet machine to yield tablets each weighing 150 mg.

Formulation 5

Capsules suitable for treating motion sickness each containing 20 mg of medicament are made as follows:

(±)-6-(5-oxadiazolyl)-4-(di-methylamino)-1,3,4,5-tetrahydrobenz[cd]indole	20 mg
Starch	169 mg
Magnesium stearate	1 mg
Total	190 mg

The active ingredient, cellulose, starch and magnesium stearate are blended, passed through a No. 20 mesh U.S. sieve, and filled into hard gelatin capsules in 190 mg quantities.

Formulation 6

Suppositories each containing 225 mg of active ingredient are made as follows:

(+)-6-(2-tetrazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole	225 mg
Saturated fatty acid glycerides to	2,000 mg

The active ingredient is passed through a No. 60 mesh U.S. sieve and suspended in the saturated fatty acid glycerides previously melted using the minimum heat necessary. The mixture is then poured into a suppository mold of nominal 2 g capacity and allowed to cool.

Formulation 7

Suspensions suitable for treating emesis each containing 50 mg of medicament per 5 ml dose are made as follows:

5	(±)-6-(2-thiazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole	50 mg
	Xanthan gum	4 mg
	Sodium carboxymethyl cellulose (11%)	
10	Microcrystalline cellulose (89%)	50 mg
	Sucrose	1.75 g
	Sodium benzoate	10 mg
15	Flavor	q.v.
	Color	q.v.
20	Purified water to	5 ml

The medicament, sucrose and xanthan gum are blended, passed through a No. 10 mesh U.S. sieve, and then mixed with a previously made solution of the microcrystalline cellulose and sodium carboxymethylcellulose in water. The sodium benzoate, flavor and color are diluted with some of the water and added with stirring. Sufficient water is then added to produce the required volume.

Formulation 8

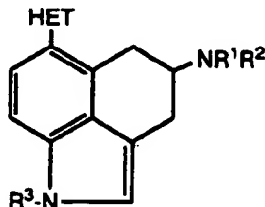
Capsules suitable for treating motion sickness each containing 50 mg of medicament are made as follows:

30	(+)-6-(5-isoxazolyl)-4-(dimethylamino)-1,3,4,5-tetrahydrobenz[cd]indole	50 mg
	Starch	507 mg
	Magnesium stearate	3 mg
35	Total	560 mg

The active ingredient, cellulose, starch and magnesium stearate are blended, passed through a No. 20 mesh U.S. sieve, and filled into hard gelatin capsules.

Claims

1. The use of a compound, or pharmaceutically acceptable salt thereof, of the formula



wherein:

R¹ is hydrogen, C₁-C₄ alkyl, C₃-C₄ alkenyl, cyclopropylmethyl, aryl(C₁-C₄ alkyl), -(CH₂)ₙS(C₁-C₄ alkyl), -C(O)R⁴, or -(CH₂)ₙC(O)NR⁵R⁶;

R² is hydrogen, C₁-C₄ alkyl, cyclopropylmethyl or C₃-C₄ alkenyl;

R³ is hydrogen, C₁-C₄ alkyl or an amino blocking group;

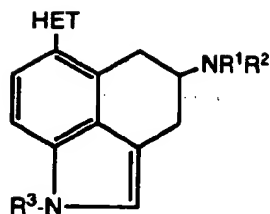
n is 1-4;

R⁴ is hydrogen, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy or phenyl;

R⁵ and R⁶ are independently hydrogen, C₁-C₄ alkyl, or C₅-C₈ cycloalkyl with the proviso that when one of R⁵ or R⁶ is a cycloalkyl the other is hydrogen;

HET is a tetrazolyl ring, a substituted tetrazolyl ring or an aromatic 5- or 6-membered heterocyclic ring, said ring having from one to three heteroatoms which are the same or different and which are selected from the group consisting of sulfur, oxygen, and nitrogen with the proviso that the 6-membered heterocyclic ring can only contain carbon and nitrogen and with the further proviso that the 5-membered ring contains no more than one oxygen or one sulfur but not both oxygen and sulfur; to prepare a medicament for treating emesis or motion sickness.

2. The use according to Claim 1 wherein the compound is one in which R¹ and R² are independently C₁-C₃ alkyl.
3. The use according to Claim 1 or Claim 2 wherein the compound is one in which R³ is hydrogen.
4. The use according to any one of Claims 1 through 3 wherein HET is an isoxazole, an oxazole, a pyrazole or an oxadiazole.
5. The use according to Claim 1 wherein the compound is selected from the group consisting of (+)(4S)-6-(3-isoxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(5-isoxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (±)-6-(3-pyrazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(4-pyrazolyl)-4-(dimethylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(4-pyridyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (±)-6-(2-pyridyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(2-thiazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(5-thiazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(2-oxadiazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(5-isoxazolyl)-4-[di(cyclopropylmethyl)amino]-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(3-isoxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(3-pyrazolyl)-4-[di(cyclopropylmethyl)amino]-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(3-pyrazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(3-phenyloxadiazol-5-yl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(2-oxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(5-oxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole; (-)(4R)-6-(5-oxazolyl)-4-[di(cyclopropylmethyl)amino]-1,3,4,5-tetrahydrobenz[cd]indole; or a pharmaceutically acceptable salt thereof.
6. The use according to Claim 5 wherein the compound is (-)(4R)-6-(2-oxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole or a pharmaceutically acceptable salt thereof.
7. The use according to Claim 5 wherein the compound is (-)(4R)-6-(5-oxazolyl)-4-(di-n-propylamino)-1,3,4,5-tetrahydrobenz[cd]indole or a pharmaceutically acceptable salt thereof.
8. The use according to Claim 5 wherein the compound is (-)(4R)-6-(5-oxazolyl)-4-[di(cyclopropylmethyl)amino]-1,3,4,5-tetrahydrobenz[cd]indole or a pharmaceutically acceptable salt thereof.
9. A pharmaceutical formulation adapted for the treatment of emesis or motion sickness comprising a compound as set forth in any one of Claims 1 through 8, or a pharmaceutically acceptable salt thereof, in combination with one or more pharmaceutically acceptable carriers, diluents or excipients therefor.
10. A compound of the formula



wherein:

R¹ is hydrogen, C₁-C₄ alkyl, C₃-C₄ alkenyl, cyclopropylmethyl, aryl(C₁-C₄ alkyl), -(CH₂)ₙS(C₁-C₄ alkyl), -C(O)R⁴, or -(CH₂)ₙC(O)NR⁵R⁶;

R² is hydrogen, C₁-C₄ alkyl, cyclopropylmethyl or C₃-C₄ alkenyl;

R³ is hydrogen, C₁-C₄ alkyl or an amino blocking group;

n is 1-4;

R⁴ is hydrogen, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy or phenyl;

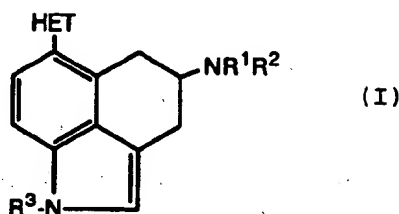
R⁵ and R⁶ are independently hydrogen, C₁-C₄ alkyl, or C₅-C₈ cycloalkyl with the proviso that when one of R⁵ or R⁶ is a cycloalkyl the other is hydrogen;

HET is a tetrazolyl ring or a substituted tetrazolyl ring; or a pharmaceutically acceptable salt thereof.

11. A pharmaceutical formulation comprising a compound of Claim 10, or a pharmaceutically acceptable salt thereof, in combination with one or more pharmaceutically acceptable carriers, diluents or excipients thereof.

12. A compound of Claim 10 for use in treating serotonin related disorders.

13. A process for preparing a compound of Formula (I)



wherein:

R¹ is hydrogen, C₁-C₄ alkyl, C₃-C₄ alkenyl, cyclopropylmethyl, aryl(C₁-C₄ alkyl), -(CH₂)ₙS(C₁-C₄ alkyl), -C(O)R⁴, or -(CH₂)ₙC(O)NR⁵R⁶;

R² is hydrogen, C₁-C₄ alkyl, cyclopropylmethyl or C₃-C₄ alkenyl;

R³ is hydrogen, C₁-C₄ alkyl or an amino blocking group;

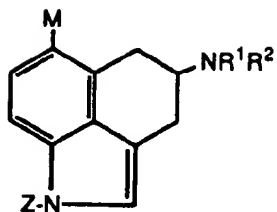
n is 1-4;

R⁴ is hydrogen, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy or phenyl;

R⁵ and R⁶ are independently hydrogen, C₁-C₄ alkyl, or C₅-C₈ cycloalkyl with the proviso that when one of R⁵ or R⁶ is a cycloalkyl the other is hydrogen;

HET is a tetrazolyl ring or a substituted tetrazolyl ring; or a pharmaceutically acceptable salt thereof, which comprises;

a) reacting a 4-amino-6-metallo-substituted tetrahydrobenz[cd]indole of the formula



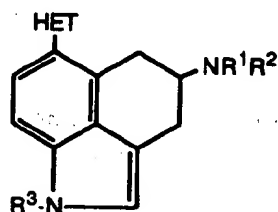
wherein R¹ and R² are as set forth above; Z is an amino protecting group and M is a metallo moiety, with a heterocyclic compound of the formula

HET-L,

wherein HET is as defined above and L is a leaving group, so as to provide a compound of Formula I

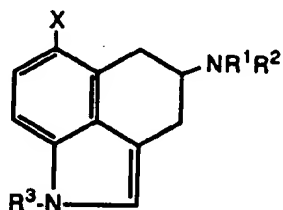
wherein R³ is an amino protecting group;

b) deprotecting a compound of the formula



wherein HET, R¹ and R² are as defined above and R³ is an amino protecting group so as to provide a compound of Formula I wherein R³ is hydrogen;

c) reacting a 4-amino-6-halo-substituted tetrahydrobenz[cd]indole of the formula

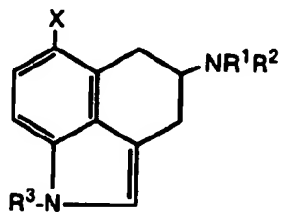


wherein R¹, R², and R³ are as defined above and X is halo with an organometallic derivative of the formula

M-HET

where HET is as defined above and M is lithium, magnesium, zinc, tin, mercury or boronic acid;

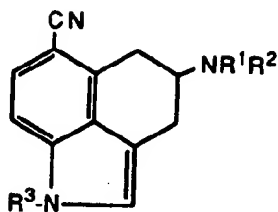
d) reacting a 4-amino-6-halo-substituted tetrahydrobenz[cd]indole of the formula



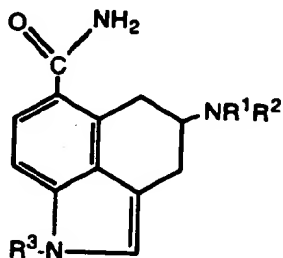
wherein R¹, R² and R³ are as defined above and X is halo with a compound of the formula

H-HET

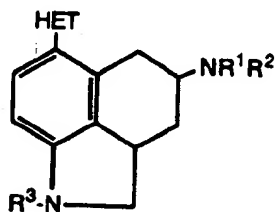
where HET is as defined above and H is hydrogen, in the presence of a catalyst;
e) reacting a nitrile of the formula



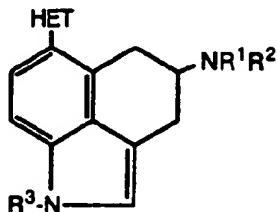
wherein R¹, R² and R³ are as defined above, with an azide so as to provide a compound wherein HET is a tetrazolyl ring;
f) reacting a compound of the formula



wherein R¹, R² and R³ are as defined above, with an azide so as to provide a compound wherein HET is a tetrazolyl ring;
g) oxidizing a compound of the formula



wherein HET, R¹, R² and R³ are as defined above, so as to provide a compound of Formula I; or
h) reacting a compound of the formula



wherein HET, R¹, R² and R³ are as defined above, with a pharmaceutically acceptable organic or inorganic acid so as to form a pharmaceutically acceptable acid addition salt of such compound.



European Patent
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EUROPEAN SEARCH REPORT

Application Number

EP 93 30 7747

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
X	CA-A-2 064 384 (ELI LILLY & COMPANY) 29 September 1992 * the whole document *	1-13	C07D403/04 A61K31/40
P,X	& EP-A-0 506 363 30 September 1992 ---	1-13	
Y	WO-A-9 004 396 (FARMITALIA CARLO ERBA S.R.L.) 3 May 1990 * the whole document *	1-13	
D,Y	JOURNAL OF MEDICINAL CHEMISTRY vol. 31, no. 9, September 1988, pages 1746 - 1753 FLAUGH M.E. ET AL. '6-Substituted 1,3,4,5-tetrahydrobenz[cd]indol-4-amines: potent serotonin agonists' * the whole document *	1-13	
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
			C07D A61K
The present search report has been drawn up for all claims			
Place of search MUNICH		Date of completion of the search 07 DECEMBER 1993	Examiner HARTRAMPF G.W.
CATEGORY OF CITED DOCUMENTS		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			

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